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(54) Title: BORRELIA BURGDORFERI POLYNUCLEOTIDES AND SEQUENCES

(57) Abstract

The present invention provides polynucleotide sequences of the genome of Borrelia Burgdorferi, polypeptide sequences encoded by the polynucleotide sequences, corresponding polynucleotides and polypeptides, vectors and hosts comprising the polynucleotides, and assays and other uses thereof. The present invention further provides polynucleotide and polypeptide sequence information stored on computer readable media, and computer-based systems and methods which facilitate its use.

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Borrelia burgdorferi Polynucleotides and Sequences

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Field of the Invention

The present invention relates to the field of molecular biology. In particular, it relates to, among other things, nucleotide sequences of *Borrelia burgdorferi*, contigs, ORFs, fragments, probes, primers and related polynucleotides thereof, peptides and polypeptides encoded by the sequences, and uses of the polynucleotides and sequences thereof, such as in fermentation, polypeptide production, assays and pharmaceutical development, among others.

Statement as to Rights to Inventions Made Under Federally-Sponsored Research and Development

Part of the work performed during development of this invention utilized U.S. Government funds. The U.S. Government may have certain rights in the invention - DE-FC02-95ER61962; DE-FC02-95ER61963; and NAGW 2554.

Background of the Invention

Spirochetes are a family of motile, unicellular, spiral-shaped bacteria which share a number of structural characteristics. Three genera of the spirochetes are pathogenic in humans: (a) *Treponema*, which includes the pathogens that cause syphilis (*T. pallidum*), yaws (*T. pertenue*), and pinta (*T. carateum*); (b) *Borrelia*, which includes the pathogens that cause epidemic and endemic relapsing fever and Lyme disease; and (c) *Leptospira*, which includes a wide variety of small spirochetes that cause mild to serious systemic human illness (Koff, A. B. and Rosen, T. *J. Am. Acad. Dermatol.* **29:**519-535 (1993)).

Lyme borreliosis, more commonly known as Lyme disease, is presently the most common human disease in the United States transmitted by an arthropod vector. Centers for Disease Control, Morbid. Mortal. Weekly Rep. 44:590-591 (1995). Further, infection of household pets, such as dogs, is a considerable problem. The causative agent of this affliction is the spirochete *Borrelia burgdorferi*, which is generally transmitted to mammalian hosts by feeding ticks. Barbour, A. and Fish, D. Science 260:1610-1616 (1993). Once the bacteria pass through the skin they disseminate and produce a variety of clinical manifestations. Diagnosis of this disease is often made serologically by the identification of antiborrelial antibodies. Hilton, E. et al., J. Clin. Microbiol. 35:774-776 (1997).

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While initial symptoms often include a rash at the infection point, Lyme disease is a multisystemic disorder that may include arthritic, carditic, and neurological manifestations. While antibiotics are currently used to treat active cases of Lyme disease, *B. burgdorferi* appears to be able to persist even after prolonged antibiotic treatment. Further, *B. burgdorferi* can persist for years in a mammalian host even in the presence of an active immune response. Straubinger, R. et al., J. Clin. Microbiol. 35:111-116 (1997); Steere, A., N. Engl. J. Med. 321:586-596 (1989).

Animal models have proven useful for studying the progression of Lyme disease, methods for preventing this disease, and immunological responses to antigenic challenges with *B. burgdorferi* proteins. Garcia-Monoco, J. et al., J. Infect. Dis. 175:1243-1245 (1997). Using a canine model, Starubinger, R. et al., Infect. Immun. 65:1273-1285 (1977), demonstrated that *B. burgdorferi* migrates into joints and induces up-regulation of interleukin-8 in synovial membranes. Similarly, *B. burgdorferi* induction of interleukin-8 production has been demonstrated in cultured human endothelial cells. Burns, M. et al., Infect. Immun. 65:1217-1222 (1997).

Antigenic heterogeneity has been postulated as a mechanism used by *B. burgdorferi* for evasion of host immune responses. Schwan, T. et al., Can. J. Microbiol. 37:450-454 (1991). In support of this mechanism, antigenic variation has been described with other pathogenic bacteria. Hagbloom, P. et al., Nature 315:156-158 (1985). Further, cassette type genetic recombination of genes encoding *B. burgdorferi* surface proteins has been shown to decrease the antigenicity of these organisms to antibodies generated against strains which have not undone the same recombination. Zhang, J. et al., Cell 89:275-285 (1997).

A number of different types of Lyme disease vaccines have been tested and shown to induce immunological responses. Whole-cell *B. burgdorferi* vaccines have been shown to induce both immunological responses and protective immunity in several animal models. Reviewed in Wormser, G., Clin. Infect. Dis. 21:1267-1274 (1995). For example, dogs inoculated with a chemically inactivated whole-cell vaccine primarily develop antibodies to outer surface membrane proteins of the administered organism. Further, passive immunity has been also demonstrated in animals using *B. burgdorferi* specific antisera. Similarly, passive immunity is conferred human by the administration of sera obtained from Lyme disease patients.

While whole-cell Lyme disease vaccines confer protective immunity in animal models, use of such vaccines presents the risk that responsive antibodies will be generated which cross react with human antigens. Reviewed in Wormser, G., supra. This problem is at least partly the result of the production of *B. burgdorferi* specific antibodies which cross-react with hepatocytes and both muscle and nerve cells. *B. burgdorferi* heat shock proteins and the 41-kd flagellin subunit are believed to contain the antigens against which these cross-reactive antibodies are generated.

It is clear that the etiology of diseases mediated or exacerbated by *B. burgdorferi* genes, and that characterizing the genes and their patterns of expression would add dramatically to our

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understanding of the organism and its host interactions. Knowledge of *B. burgdorferi* genes and genomic organization would dramatically improve understanding of disease etiology and lead to improved and new ways of preventing, ameliorating, arresting and reversing diseases. Moreover, characterized genes and genomic fragments of *B. burgdorferi* would provide reagents for, among other things, detecting, characterizing and controlling *B. burgdorferi* infections. There is a need therefore to characterize the genome of *B. burgdorferi* and for polynucleotides and sequences of this organism.

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SUMMARY OF THE INVENTION

The present invention is based on the sequencing of fragments of the *Borrelia burgdorferi* genome. The primary nucleotide sequences which were generated are provided in SEQ ID NOS:1-155.

The present invention provides the complete nucleotide sequence of the *Borrelia burgdorferi* chromosome and 154 contigs representing the majority of the sequence of the B. burgdorferi extrachromosomal elements, all of which are listed in tables below and set out in the Sequence Listing submitted herewith, and representative fragments thereof, in a form which can be readily used, analyzed, and interpreted by a skilled artisan. In one embodiment, the present invention is provided as contiguous strings of primary sequence information corresponding to the nucleotide sequences depicted in SEQ ID NOS: 1-155.

The present invention further provides nucleotide sequences which are at least 95%, 96%, 97%, 98%, and 99%, identical to the nucleotide sequences of SEQ ID NOS:1-155, ORF IDs and corresponding ORFs.

The nucleotide sequences of SEQ ID NOS:1-155, ORF ID or ORF within, a representative fragment thereof, or a nucleotide sequence which is at least 95% identical to said nucleotide sequence may be provided in a variety of mediums to facilitate its use. In one application of this embodiment, the sequences of the present invention are recorded on computer readable media. Such media includes, but is not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD-ROM; electrical storage media such as RAM and ROM; and hybrids of these categories such as magnetic/optical storage media.

The present invention further provides systems, particularly computer-based systems which contain the sequence information herein described stored in a data storage means. Such systems are designed to identify commercially important fragments of the *Borrelia burgdorferi* genome.

Another embodiment of the present invention is directed to fragments of the *Borrelia* burgdorferi genome having particular structural or functional attributes. Such fragments of the *Borrelia burgdorferi* genome of the present invention include, but are not limited to, fragments which encode peptides, hereinafter referred to as open reading frames or ORFs, fragments which modulate the expression of an operably linked ORF, hereinafter referred to as expression

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modulating fragments or EMFs, and fragments which can be used to diagnose the presence of *Borrelia burgdorferi* in a sample, hereinafter referred to as diagnostic fragments or DFs.

Each of the ORF IDs and ORFs in fragments of the *Borrelia burgdorferi* genome disclosed in Tables 1-6, and the EMFs found 5' prime of the initiation codon, can be used in numerous ways as polynucleotide reagents. For instance, the sequences can be used as diagnostic probes or amplification primers for detecting or determining the presence of a specific microbe in a sample, to selectively control gene expression in a host and in the production of polypeptides, such as polypeptides encoded by ORFs of the present invention, particular those polypeptides that have a pharmacological activity.

The present invention further includes recombinant constructs comprising one or more fragments of the *Borrelia burgdorferi* genome of the present invention. The recombinant constructs of the present invention comprise vectors, such as a plasmid or viral vector, into which a fragment of the *Borrelia burgdorferi* has been inserted.

The present invention further provides host cells containing any of the isolated fragments of the *Borrelia burgdorferi* genome of the present invention. The host cells can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic cell, such as a yeast cell, or a procaryotic cell such as a bacterial cell.

The present invention is further directed to isolated polypeptides and proteins encoded by ORFs of the present invention. A variety of methods, well known to those of skill in the art, routinely may be utilized to obtain any of the polypeptides and proteins of the present invention. For instance, polypeptides and proteins of the present invention having relatively short, simple amino acid sequences readily can be synthesized using commercially available automated peptide synthesizers. Polypeptides and proteins of the present invention also may be purified from bacterial cells which naturally produce the protein. Yet another alternative is to purify polypeptide and proteins of the present invention from cells which have been altered to express them.

The invention further provides methods of obtaining homologs of the fragments of the *Borrelia burgdorferi* genome of the present invention and homologs of the proteins encoded by the ORFs of the present invention. Specifically, by using the nucleotide and amino acid sequences disclosed herein as a probe or as primers, and techniques such as PCR cloning and colony/plaque hybridization, one skilled in the art can obtain homologs.

The invention further provides antibodies which selectively bind polypeptides and proteins of the present invention. Such antibodies include both monoclonal and polyclonal antibodies.

The invention further provides hybridomas which produce the above-described antibodies. A hybridoma is an immortalized cell line which is capable of secreting a specific monoclonal antibody.

The present invention further provides methods of identifying test samples derived from cells which express one of the ORFs of the present invention, or a homolog thereof. Such

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methods comprise incubating a test sample with one or more of the antibodies of the present invention, or one or more of the DFs of the present invention, under conditions which allow a skilled artisan to determine if the sample contains the ORF or product produced therefrom.

In another embodiment of the present invention, kits are provided which contain the necessary reagents to carry out the above-described assays.

Specifically, the invention provides a compartmentalized kit to receive, in close confinement, one or more containers which comprises: (a) a first container comprising one of the antibodies, or one of the DFs of the present invention; and (b) one or more other containers comprising one or more of the following: wash reagents, reagents capable of detecting presence of bound antibodies or hybridized DFs.

Using the isolated proteins of the present invention, the present invention further provides methods of obtaining and identifying agents capable of binding to a polypeptide or protein encoded by one of the ORFs of the present invention. Specifically, such agents include, as further described below, antibodies, peptides, carbohydrates, pharmaceutical agents and the like. Such methods comprise steps of: (a)contacting an agent with an isolated protein encoded by one of the ORFs of the present invention; and (b)determining whether the agent binds to said protein.

The present genomic sequences of *Borrelia burgdorferi* will be of great value to all laboratories working with this organism and for a variety of commercial purposes. Many fragments of the *Borrelia burgdorferi* genome will be immediately identified by similarity searches against GenBank or protein databases and will be of immediate value to *Borrelia burgdorferi* researchers and for immediate commercial value for the production of proteins or to control gene expression.

The methodology and technology for elucidating extensive genomic sequences of bacterial and other genomes has and will greatly enhance the ability to analyze and understand chromosomal organization. In particular, sequenced contigs and genomes will provide the models for developing tools for the analysis of chromosome structure and function, including the ability to identify genes within large segments of genomic DNA, the structure, position, and spacing of regulatory elements, the identification of genes with potential industrial applications, and the ability to do comparative genomic and molecular phylogeny.

DESCRIPTION OF THE FIGURES

FIGURE 1 is a block diagram of a computer system (102) that can be used to implement computer-based systems of present invention.

FIGURE 2 is a schematic diagram depicting the data flow and computer programs used to collect, assemble, edit and annotate the contigs of the *Borrelia burgdorferi* genome of the present invention. Both Macintosh and Unix platforms are used to handle the AB 373 and 377 sequence data files, largely as described in Kerlavage *et al.*, *Proceedings of the Twenty-Sixth*

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Annual Hawaii International Conference on System Sciences, 585, IEEE Computer Society Press, Washington D.C. (1993). Factura (AB) is a Macintosh program designed for automatic vector sequence removal and end-trimming of sequence files. The program Loadis runs on a Macintosh platform and parses the feature data extracted from the sequence files by Factura to the Unix based Borrelia burgdorferi relational database. Assembly of contigs (and whole genome sequences) is accomplished by retrieving a specific set of sequence files and their associated features using Extrseq, a Unix utility for retrieving sequences from an SOL database. The resulting sequence file is processed to trim portions of the sequences with a high rate ambiguous nucleotides. The sequence files were assembled using TIGR Assembler, an assembly engine designed at The Institute for Genomic Research (TIGR) for rapid and accurate assembly of thousands of sequence fragments. The collection of contigs generated by the assembly step is loaded into the database with the lassie program. Identification of open reading frames (ORFs) is accomplished by processing contigs with zorf. The ORFs are searched against B. burgdorferi sequences from GenBank and against all protein sequences using the BLASTN and BLASTP programs, described in Altschul et al., J. Mol. Biol. 215: 403-410 (1990). Results of the ORF determination and similarity searching steps were loaded into the database. As described below, some results of the determination and the searches are set out in Tables 1-6.

DETAILED DESCRIPTION OF ILLUSTRATIVE EMBODIMENTS

The present invention is based on the sequencing of fragments of the *Borrelia burgdorferi* genome and analysis of the sequences. The primary nucleotide sequences generated by sequencing the fragments are provided in SEQ ID NOS: 1-155. (As used herein, the "primary sequence" refers to the nucleotide sequence represented by the IUPAC nomenclature system.) SEQ ID NOS:1-155

In addition, the present invention provides the nucleotide sequences of SEQ ID NOS: 1-155, or representative fragments thereof, in a form which can be readily used, analyzed, and interpreted by a skilled artisan.

As used herein, a "representative fragment of the nucleotide sequence depicted in SEQ ID NOS:1-155" refers to any portion of the SEQ ID NOS: 1-155 which is not presently represented within a publicly available database. Preferred representative fragments of the present invention are *Borrelia burgdorferi* open reading frames (ORFs) represented by ORF IDs, expression modulating fragments (EMFs) and diagnostic fragments (DFs)which can be used to diagnose the presence of *Borrelia burgdorferi* in sample. A non-limiting identification of preferred representative portions are provided in Tables 1-6 as ORF IDs. As discussed in detail below, the information provided in SEQ ID NOS:1-155 and in Tables 1-6 together with routine cloning, synthesis, sequencing and assay methods will enable those skilled in the art to clone and sequence all "representative fragments" of interest, including ORFs encoding a large variety of *Borrelia burgdorferi* proteins.

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The present invention is further directed to nucleic acid molecules encoding portions or fragments of the nucleotide sequences described herein. Fragments include portions of the nucleotide sequences of Table 1-6 (ORF IDs) and SEQ ID NOS:1-155, at least 10 contiguous nucleotides in length selected from any two integers, one of which representing a 5' nucleotide position and a second of which representing a 3' nucleotide position, where the first nucleotide for each nucleotide sequence in SEQ ID NOS:1-155 is position 1 (therefore, the sequence postions for each ORF ID is determined by the numbering of the SEQ ID comprising the ORF ID). That is, every combination of a 5' and 3' nucleotide position that a fragment at least 10 contiguous nucleotides in length could occupy is included in the invention. At least means a fragment may be 10 contiguous nucleotide bases in length or any integer between 10 and the length of an entire nucleotide sequence of SEQ ID NOS:1-155 minus 1. Therefore, included in the invention are contiguous fragments specified by any 5' and 3' nucleotide base positions of a nucleotide sequences of SEQ ID NOS:1-155 wherein the contiguous fragment is any integer between 10 and the length of an entire nucleotide sequence minus 1.

Further, the invention includes polynucleotides comprising fragments specified by size, in nucleotides, rather than by nucleotide positions. The invention includes any fragment size, in contiguous nucleotides, selected from integers between 10 and the length of an entire ORF ID or SEQ ID NO:, minus 1. Preferred sizes of contiguous nucleotide fragments include 20 nucleotides, 30 nucleotides, 40 nucleotides, 50 nucleotides. Other preferred sizes of contiguous nucleotide fragments, which may be useful as diagnostic probes and primers, include fragments 50-300 nucleotides in length which include, as discussed above, fragment sizes representing each integer between 50-300. Larger fragments are also useful according to the present invention corresponding to most, if not all, of the nucleotide sequences shown in Tables 1-6 (ORF IDs) and SEQ ID NOS:1-155. The preferred sizes are, of course, meant to exemplify not limit the present invention as all size fragments, representing any integer between 10 and the length of an entire nucleotide sequence minus 1, of each ORF ID and SEQ ID NO:, are included in the invention.

The present invention also provides for the exclusion of any fragment, specified by 5' and 3' base positions or by size in nucleotide bases as described above for any ORF ID or SEQ ID NOS:1-155. Any number of fragments of nucleotide sequences in ORF IDs or SEQ ID NOS:1-155, specified by 5' and 3' base positions or by size in nucleotides, as described above, may be excluded from the present invention.

While the presently disclosed sequences of SEQ ID NOS: 1-155 are highly accurate, sequencing techniques are not perfect and, in relatively rare instances, further investigation of a fragment or sequence of the invention may reveal a nucleotide sequence error present in a nucleotide sequence disclosed in SEQ ID NOS: 1-155. However, once the present invention is made available (*i.e.*, once the information in SEQ ID NOS: 1-155 and Tables 1-6 has been made available), resolving a rare sequencing error in SEQ ID NOS: 1-155 will be well within the skill

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of the art. The present disclosure makes available sufficient sequence information to allow any of the described contigs or portions thereof to be obtained readily by straightforward application of routine techniques. Further sequencing of such polynucleotide may proceed in like manner using manual and automated sequencing methods which are employed ubiquitous in the art. Nucleotide sequence editing software is publicly available. For example, Applied Biosystem's (AB) AutoAssembler can be used as an aid during visual inspection of nucleotide sequences. By employing such routine techniques potential errors readily may be identified and the correct sequence then may be ascertained by targeting further sequencing effort, also of a routine nature, to the region containing the potential error.

Even if all of the very rare sequencing errors in SEQ ID NOS: 1-155 were corrected, the resulting nucleotide sequences would still be at least 95% identical, nearly all would be at least 99% identical, and the great majority would be at least 99.9% identical to the nucleotide sequences of SEQ ID NOS: 1-155.

As discussed elsewhere herein, polynucleotides of the present invention readily may be obtained by routine application of well known and standard procedures for cloning and sequencing DNA. Detailed methods for obtaining libraries and for sequencing are provided below, for instance. A wide variety of *Borrelia burgdorferi* strains that can be used to prepare *B. burgdorferi* genomic DNA for cloning and for obtaining polynucleotides of the present invention are available to the public from recognized depository institutions, such as the American Type Culture Collection (ATCC). While the present invention is enabled by the sequences and other information herein disclosed, the *B. burgdorferi* strain that provided the DNA of the present Sequence Listing, has been deposited with the ATCC, 10801 University Blvd. Manassas, VA 20110-2209, as Deposit No. 202012, on 8 August 1997. The ATCC Deposit is provided merely as a convenience to those of skill in the art. Reference to the deposit is not a waiver of any rights of the inventors or their assignees in the present subject matter.

The nucleotide sequences of the genomes from different strains of *Borrelia burgdorferi* differ somewhat. However, the nucleotide sequences of the genomes of all *Borrelia burgdorferi* strains will be at least 95% identical, in corresponding part, to the nucleotide sequences provided in SEQ ID NOS: 1-155 and the ORF IDs within. Nearly all will be at least 99% identical and the great majority will be 99.9% identical.

The present application is further directed to nucleic acid molecules at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleic acid sequence shown in SEQ ID NOS: 1-155 and the ORF IDs within. The above nucleic acid sequences are included irrespective of whether they encode a polypeptide having *B. burgdorferi* activity. This is because even where a particular nucleic acid molecule does not encode a polypeptide having *B. burgdorferi* activity, one of skill in the art would still know how to use the nucleic acid molecule, for instance, as a hybridization probe. Uses of the nucleic acid molecules of the present invention that do not encode a polypeptide having *B. burgdorferi* activity include, *inter alia*, isolating a *B. burgdorferi* gene or allelic variants thereof from a DNA library, and detecting *B. burgdorferi* mRNA expression from

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biological or environmental samples, suspected of containing *B. burgdorferi* by Northern Blot, PCR, or similar analysis.

Preferred, are nucleic acid molecules having sequences at least 90%, 95%, 96%, 97%, 98% or 99% identical to the nucleic acid sequence shown in SEQ ID NOS: 1-155, the ORF IDs, and the ORF within each ORF ID, which do, in fact, encode a polypeptide having *B. burgdorferi* protein activity. By "a polypeptide having *B. burgdorferi* activity" is intended polypeptides exhibiting activity similar, but not necessarily identical, to an activity of the *B. burgdorferi* protein of the invention, as measured in a particular biological assay suitable for measuring activity of the specified protein.

Due to the degeneracy of the genetic code, one of ordinary skill in the art will immediately recognize that a large number of the nucleic acid molecules having a sequence at least 90%, 95%, 96%, 97%, 98%, or 99% identical to the nucleic acid sequences shown in SEQ ID NOS: 1-155, the ORF IDs, and the ORF within each ORF ID, will encode a polypeptide having *B. burgdorferi* protein activity. In fact, since degenerate variants of these nucleotide sequences all encode the same polypeptide, this will be clear to the skilled artisan even without performing the above described comparison assay. It will be further recognized in the art that, for such nucleic acid molecules that are not degenerate variants, a reasonable number will also encode a polypeptide having *B. burgdorferi* protein activity. This is because the skilled artisan is fully aware of amino acid substitutions that are either less likely or not likely to significantly effect protein function (e.g., replacing one aliphatic amino acid with a second aliphatic amino acid), as further described below.

The biological activity or function of the polypeptides of the present invention are expected to be similar or identical to polypeptides from other bacteria that share a high degree of structural identity/similarity. Tables 1, 2, 4, and 5 lists accession numbers and descriptions for the closest matching sequences of polypeptides available through Genbank. It is therefore expected that the biological activity or function of the polypeptides of the present invention will be similar or identical to those polypeptides from other bacterial genuses, species, or strains listed in Tables 1, 2, 4, and 5.

By a polynucleotide having a nucleotide sequence at least, for example, 95% "identical" to a reference nucleotide sequence of the present invention, it is intended that the nucleotide sequence of the polynucleotide is identical to the reference sequence except that the polynucleotide sequence may include up to five point mutations per each 100 nucleotides of the reference nucleotide sequence encoding the *B. burgdorferi* polypeptide. In other words, to obtain a polynucleotide having a nucleotide sequence at least 95% identical to a reference nucleotide sequence, up to 5% of the nucleotides in the reference sequence may be deleted, inserted, or substituted with another nucleotide. The query sequence may be an entire sequence shown in SEQ ID NOS: 1-155, an ORF ID, or the ORF within each ORF ID, or any fragment specified as described herein.

As a practical matter, whether any particular nucleic acid molecule or polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to a nucleotide sequence of the presence invention can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al. See Brutlag et al. (1990) Comp. App. Biosci. 6:237-245. In a sequence alignment the query and subject sequences are both DNA sequences. An RNA sequence can be compared by first converting U's to T's. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB alignment of DNA sequences to calculate percent identity are: Matrix=Unitary, k-tuple=4, Mismatch Penalty=1, Joining Penalty=30, Randomization Group Length=0, Cutoff Score=1, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject nucleotide sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence because of 5' or 3' deletions, not because of internal deletions, a manual correction must be made to the results. This is because the FASTDB program does not account for 5' and 3' truncations of the subject sequence when calculating percent identity. For subject sequences truncated at the 5' or 3' ends, relative to the query sequence, the percent identity is corrected by calculating the number of bases of the query sequence that are 5' and 3' of the subject sequence, which are not matched/aligned, as a percent of the total bases of the query sequence. Whether a nucleotide is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This corrected score is what is used for the purposes of the present invention. Only nucleotides outside the 5' and 3' nucleotides of the subject sequence, as displayed by the FASTDB alignment, which are not matched/aligned with the query sequence, are calculated for the purposes of manually adjusting the percent identity score.

For example, a 90 nucleotide subject sequence is aligned to a 100 nucleotide query sequence to determine percent identity. The deletions occur at the 5' end of the subject sequence and therefore, the FASTDB alignment does not show a matched/alignment of the first 10 nucleotides at 5' end. The 10 unpaired nucleotides represent 10% of the sequence (number of nucleotides at the 5' and 3' ends not matched/total number of nucleotides in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 nucleotides were perfectly matched the final percent identity would be 90%. In another example, a 90 nucleotide subject sequence is compared with a 100 nucleotide query sequence. This time the deletions are internal deletions so that there are no nucleotides on the 5' or 3' of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only nucleotides 5' and 3' of the

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subject sequence which are not matched/aligned with the query sequence are manually corrected for. No other manual corrections are to made for the purposes of the present invention.

COMPUTER RELATED EMBODIMENTS

The nucleotide sequences provided in SEQ ID NOS: 1-155, including ORF IDs and corresponding ORFs, a representative fragment thereof, or a nucleotide sequence at least 95%, preferably at least 96%, 97%, 98% or 99%, and most preferably at least 99.9% identical to said nucleotide sequences may be "provided" in a variety of mediums to facilitate use thereof. As used herein, provided refers to a manufacture, other than an isolated nucleic acid molecule, which contains a nucleotide sequence of the present invention, a representative fragment thereof, or a nucleotide sequence at least 95%, preferably at least 99% and most preferably at least 99.9% identical to a polynucleotide of the present invention. Such a manufacture provides a large portion of the *Borrelia burgdorferi* genome and parts thereof (*e.g.*, a *Borrelia burgdorferi* open reading frame (ORF)) in a form which allows a skilled artisan to examine the manufacture using means not directly applicable to examining the *Borrelia burgdorferi* genome or a subset thereof as it exists in nature or in purified form.

In one application of this embodiment, a nucleotide sequence of the present invention can be recorded on computer readable media. As used herein, "computer readable media" refers to any medium which can be read and accessed directly by a computer. Such media include, but are not limited to: magnetic storage media, such as floppy discs, hard disc storage medium, and magnetic tape; optical storage media such as CD- ROM; electrical storage media such as RAM and ROM; and hybrids of these categories, such as magnetic/optical storage media. A skilled artisan can readily appreciate how any of the presently known computer readable mediums can be used to create a manufacture comprising computer readable medium having recorded thereon a nucleotide sequence of the present invention. Likewise, it will be clear to those of skill how additional computer readable media that may be developed also can be used to create analogous manufactures having recorded thereon a nucleotide sequence of the present invention.

As used herein, "recorded" refers to a process for storing information on computer readable medium. A skilled artisan can readily adopt any of the presently know methods for recording information on computer readable medium to generate manufactures comprising the nucleotide sequence information of the present invention.

A variety of data storage structures are available to a skilled artisan for creating a computer readable medium having recorded thereon a nucleotide sequence of the present invention. The choice of the data storage structure will generally be based on the means chosen to access the stored information. In addition, a variety of data processor programs and formats can be used to store the nucleotide sequence information of the present invention on computer readable medium. The sequence information can be represented in a word processing text file, formatted in commercially- available software such as WordPerfect and MicroSoft Word, or represented in the form of an ASCII file, stored in a database application, such as DB2, Sybase,

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Oracle, or the like. A skilled artisan can readily adapt any number of data-processor structuring formats (e.g., text file or database) in order to obtain computer readable medium having recorded thereon the nucleotide sequence information of the present invention.

Computer software is publicly available which allows a skilled artisan to access sequence information provided in a computer readable medium. Thus, by providing in computer readable form the nucleotide sequences of the present invention (e.g. SEQ ID NOS: 1-155), a representative fragment thereof, or a nucleotide sequence at least 95%, preferably at least 96%, 97%, 98%, 99% and most preferably at least 99.9% identical to a sequence of the present invention (e.g. SEQ ID NOS: 1-155) enables the skilled artisan routinely to access the provided sequence information for a wide variety of purposes.

The examples which follow demonstrate how software which implements the BLAST (Altschul et al., J. Mol. Biol. 215:403-410 (1990)) and BLAZE (Brutlag et al., Comp. Chem. 17:203-207 (1993)) search algorithms on a Sybase system was used to identify open reading frames (ORFs) within the Borrelia burgdorferi genome which contain homology to ORFs or proteins from both Borrelia burgdorferi and from other organisms. Among the ORFs discussed herein are protein encoding fragments of the Borrelia burgdorferi genome useful in producing commercially important proteins, such as enzymes used in fermentation reactions and in the production of commercially useful metabolites.

The present invention further provides systems, particularly computer-based systems, which contain the sequence information described herein. Such systems are designed to identify, among other things, commercially important fragments of the *Borrelia burgdorferi* genome.

As used herein, "a computer-based system" refers to the hardware means, software means, and data storage means used to analyze the nucleotide sequence information of the present invention. The minimum hardware means of the computer-based systems of the present invention comprises a central processing unit (CPU), input means, output means, and data storage means. A skilled artisan can readily appreciate that any one of the currently available computer-based system are suitable for use in the present invention.

As stated above, the computer-based systems of the present invention comprise a data storage means having stored therein a nucleotide sequence of the present invention and the necessary hardware means and software means for supporting and implementing a search means.

As used herein, "data storage means" refers to memory which can store nucleotide sequence information of the present invention, or a memory access means which can access manufactures having recorded thereon the nucleotide sequence information of the present invention.

As used herein, "search means" refers to one or more programs which are implemented on the computer-based system to compare a target sequence or target structural motif with the sequence information stored within the data storage means. Search means are used to identify fragments or regions of the present genomic sequences which match a particular target sequence or target motif. A variety of known algorithms are disclosed publicly and a variety of

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commercially available software for conducting search means are and can be used in the computer-based systems of the present invention. Examples of such software includes, but is not limited to, MacPattern (EMBL), BLASTN and BLASTX (NCBIA). A skilled artisan can readily recognize that any one of the available algorithms or implementing software packages for conducting homology searches can be adapted for use in the present computer-based systems.

As used herein, a "target sequence" can be any DNA or amino acid sequence of six or more nucleotides or two or more amino acids. A skilled artisan can readily recognize that the longer a target sequence is, the less likely a target sequence will be present as a random occurrence in the database. The most preferred sequence length of a target sequence is from about 10 to 100 amino acids or from about 30 to 300 nucleotide residues. However, it is well recognized that searches for commercially important fragments, such as sequence fragments involved in gene expression and protein processing, may be of shorter length.

As used herein, "a target structural motif," or "target motif," refers to any rationally selected sequence or combination of sequences in which the sequence(s) are chosen based on a three-dimensional configuration which is formed upon the folding of the target motif. There are a variety of target motifs known in the art. Protein target motifs include, but are not limited to, enzymic active sites and signal sequences. Nucleic acid target motifs include, but are not limited to, promoter sequences, hairpin structures and inducible expression elements (protein binding sequences).

A variety of structural formats for the input and output means can be used to input and output the information in the computer-based systems of the present invention. A preferred format for an output means ranks fragments of the *Borrelia burgdorferi* genomic sequences possessing varying degrees of homology to the target sequence or target motif. Such presentation provides a skilled artisan with a ranking of sequences which contain various amounts of the target sequence or target motif and identifies the degree of homology contained in the identified fragment.

A variety of comparing means can be used to compare a target sequence or target motif with the data storage means to identify sequence fragments of the *Borrelia burgdorferi* genome. In the present examples, implementing software which implement the BLAST and BLAZE algorithms, described in Altschul *et al.*, *J. Mol. Biol. 215:* 403-410 (1990), is used to identify open reading frames within the *Borrelia burgdorferi* genome. A skilled artisan can readily recognize that any one of the publicly available homology search programs can be used as the search means for the computer-based systems of the present invention. Of course, suitable proprietary systems that may be known to those of skill also may be employed in this regard.

Figure 1 provides a block diagram of a computer system illustrative of embodiments of this aspect of present invention. The computer system 102 includes a processor 106 connected to a bus 104. Also connected to the bus 104 are a main memory 108 (preferably implemented as random access memory, RAM) and a variety of secondary storage devices 110, such as a hard drive 112 and a removable medium storage device 114. The removable medium storage device

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114 may represent, for example, a floppy disk drive, a CD-ROM drive, a magnetic tape drive, etc. A removable storage medium 116 (such as a floppy disk, a compact disk, a magnetic tape, etc.) containing control logic and/or data recorded therein may be inserted into the removable medium storage device 114. The computer system 102 includes appropriate software for reading the control logic and/or the data from the removable medium storage device 114, once it is inserted into the removable medium storage device 114.

A nucleotide sequence of the present invention may be stored in a well known manner in the main memory 108, any of the secondary storage devices 110, and/or a removable storage medium 116. During execution, software for accessing and processing the genomic sequence (such as search tools, comparing tools, *etc.*) reside in main memory 108, in accordance with the requirements and operating parameters of the operating system, the hardware system and the software program or programs.

BIOCHEMICAL EMBODIMENTS

Other embodiments of the present invention are directed to isolated fragments of the *Borrelia burgdorferi* genome. The fragments of the *Borrelia burgdorferi* genome of the present invention include, but are not limited to fragments which encode peptides, hereinafter open reading frames (ORFs), fragments which modulate the expression of an operably linked ORF, hereinafter expression modulating fragments (EMFs) and fragments which can be used to diagnose the presence of *Borrelia burgdorferi* in a sample, hereinafter diagnostic fragments (DFs).

As used herein, an "isolated nucleic acid molecule" or an "isolated fragment of the *Borrelia burgdorferi* genome" refers to a nucleic acid molecule possessing a specific nucleotide sequence which has been subjected to purification means to reduce, from the composition, the number of compounds which are normally associated with the composition. Particularly, the term refers to the nucleic acid molecules having the sequences set out in SEQ ID NOS: 1-155, to representative fragments thereof as described above including ORF IDs and ORFs, to polynucleotides at least 95%, preferably at least 96%, 97%, 98%, or 99% and especially preferably at least 99.9% identical in sequence thereto, also as set out above.

A variety of purification means can be used to generate the isolated fragments of the present invention. These include, but are not limited to methods which separate constituents of a solution based on charge, solubility, or size.

In one embodiment, *Borrelia burgdorferi* DNA can be enzymatically sheared to produce fragments of 15-20 kb in length. These fragments can then be used to generate a *Borrelia burgdorferi* library by inserting them into lambda clones as described in the Examples below. Primers flanking, for example, an ORF, such as those enumerated in Tables 1-6 can then be generated using nucleotide sequence information provided in SEQ ID NOS: 1-155. Well known and routine techniques of PCR cloning then can be used to isolate the ORF from the lambda DNA library or *Borrelia burgdorferi* genomic DNA. Thus, given the availability of SEQ ID NOS:1-

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155, the information in Tables 1-6, and the information that may be obtained readily by analysis of the sequences of SEQ ID NOS:1-155 using methods set out above, those of skill will be enabled by the present disclosure to isolate any ORF-containing or other nucleic acid fragment of the present invention.

The isolated nucleic acid molecules of the present invention include, but are not limited to single stranded and double stranded DNA, and single stranded RNA. For purposes of numbering and reference to polynucleotide and polypeptide sequences the entire sequence of each sequence of SEQ ID NOS:1-155 is included with the first nucleotide being position 1. Therefore, for reference purposes the numbering used in the present invention is that provided in the sequence listing for SEQ ID NOS:1-155.

As used herein, an open reading frame (ORF), means a series of nucleotide triplets coding for amino acid residues without any termination codons and is a sequence translatable into protein. Further, unless specified, the term "ORF" for each ORF ID is defined by the termination codon at the 3' end and the 5' most methionine codon, at the 5' end, in frame with said 3' termination codon. Unless specified, the term "ORF" also refers to a particular polypeptide sequence defined by the ORF polynucleotide sequence, wherein the N-terminus is defined by the 5' most methionine codon in frame with the termination codon at the 3' end of the ORF ID and the C-terminus is defined by the last codon before the said 3' termination codon. As used herein, an ORF ID represents a sequence without any internal termination codons flanked by termination codons.

Tables 1-6 list ORF IDs in the *Borrelia burgdorferi* genomic contigs of the present invention that were identified as putative coding regions by the GeneMark software using organism-specific second-order Markov probability transition matrices. It will be appreciated that other criteria can be used, in accordance with well known analytical methods, such as those discussed herein, to generate more inclusive, more restrictive, or more selective lists.

The *B. burgdorferi* genome consists of one large linear chromosome containing approximately two thirds of its genetic material and multiple extrachromosomal elements (approximately 15) containing the remaining one third of its genetic material. SEQ ID NO:1 (Contig ID 1) is the complete sequence of the large linear *B. burgdorferi* chromosome. SEQ ID NOS:2-155 (Contig ID 2-155 respectively) are fragments (contigs) of the extrachromosomal elements. Tables 1-3 below relate only to SEQ ID NO:1. Tables 4-6 relate to the extrachromosomal elements (SEQ ID NOS:2-155).

Table 1 sets out ORF IDs in the *Borrelia burgdorferi* chromosome of the present invention that cover a continuous region of at least 50 bases are 95% or more identical (by BLAST analysis using default parameters) to a nucleotide sequence available through GenBank in July, 1997.

Table 2 sets out ORF IDs in the *Borrelia burgdorferi* chromosome of the present invention that are not in Table 1 and match, with a BLASTP probability score of 0.01 or less, a polypeptide sequence available through GenBank in July, 1997.

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Table 3 sets out ORF IDs in the *Borrelia burgdorferi* chromosome of the present invention that do not match significantly, by BLASTP analysis, a polypeptide sequence available through GenBank in July, 1997.

Table 4 sets out ORF IDs in the *Borrelia burgdorferi* extrachromosomal element contigs of the present invention that over a continuous region of at least 50 bases are 95% or more identical (by BLAST analysis) to a nucleotide sequence available through GenBank in July, 1997.

Table 5 sets out ORF IDs in the *Borrelia burgdorferi* extrachromosomal element contigs of the present invention that are not in Table 1 and match, with a BLASTP probability score of 0.01 or less, a polypeptide sequence available through GenBank in July, 1997.

Table 6 sets out ORF IDs in the *Borrelia burgdorferi* extrachromosomal element contigs of the present invention that do not match significantly, by BLASTP analysis, a polypeptide sequence available through GenBank in July, 1997.

In each table, the first and second columns identify the ORF ID by, respectively, contig number and ORF ID number within the contig; the third column indicates the first nucleotide of the ORF ID, counting from the 5' end of the contig strand; and the fourth column indicates the last nucleotide of the ORF ID, counting from the 5' end of the contig strand.

In Tables 1, 2, 4 and 5, column five, lists the Reference for the closest matching sequence available through GenBank. These reference numbers are the database accession numbers commonly used by those of skill in the art, who will be familiar with their denominators. Descriptions of the nomenclature are available from the National Center for Biotechnology Information. Column seven provides the BLAST identity score from the comparison of the ORF ID and the homologous gene; and column nine indicates the length in nucleotides of the highest scoring segment pair identified by the BLAST identity analysis.

The concepts of percent identity and percent similarity of two polypeptide sequences is well understood in the art. For example, two polypeptides 10 amino acids in length which differ at three amino acid positions (e.g., at positions 1, 3 and 5) are said to have a percent identity of 70%. However, the same two polypeptides would be deemed to have a percent similarity of 80% if, for example at position 5, the amino acids moieties, although not identical, were "similar" (i.e., possessed similar biochemical characteristics). As is known in the art, substitution of one amino acid for a "similar" amino acid is a conservative substitution. Generally, proteins are highly tolerant of conservative substitutions. Many programs for analysis of nucleotide or amino acid sequence similarity, such as fasta and BLAST specifically list percent identity of a matching region as an output parameter. Thus, for instance, Tables 1, 2, 4 and 5 herein enumerate the percent identity and similarity of the highest scoring segment pair in each ORF and its listed relative. Further details concerning the algorithms and criteria used for homology searches are provided below and are described in the pertinent literature highlighted by the citations provided below.

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It will be appreciated that other criteria can be used to generate more inclusive and more exclusive listings of the types set out in the tables. As those of skill will appreciate, narrow and broad searches both are useful. Thus, a skilled artisan can readily identify ORFs in contigs of the *Borrelia burgdorferi* genome other than those listed in Tables 1-6, such as ORFs which are overlapping or encoded by the opposite strand of an identified ORF in addition to those ascertainable using the computer-based systems of the present invention.

As used herein, an "expression modulating fragment," EMF, means a series of nucleotide molecules which modulates the expression of an operably linked ORF or EMF.

As used herein, a sequence is said to "modulate the expression of an operably linked sequence" when the expression of the sequence is altered by the presence of the EMF. EMFs include, but are not limited to, promoters, and promoter modulating sequences (inducible elements). One class of EMFs are fragments which induce the expression or an operably linked ORF in response to a specific regulatory factor or physiological event.

EMF sequences can be identified within the contigs of the *Borrelia burgdorferi* genome by their proximity to the ORFs provided in Tables 1-6. An intergenic segment, or a fragment of the intergenic segment, from about 10 to 200 nucleotides in length, taken from any one of the ORFs of Tables 1-6 will modulate the expression of an operably linked ORF in a fashion similar to that found with the naturally linked ORF sequence. As used herein, an "intergenic segment" refers to fragments of the *Borrelia burgdorferi* genome which are between two ORF(s) herein described. EMFs also can be identified using known EMFs as a target sequence or target motif in the computer-based systems of the present invention. Further, the two methods can be combined and used together.

The presence and activity of an EMF can be confirmed using an EMF trap vector. An EMF trap vector contains a cloning site linked to a marker sequence. A marker sequence encodes an identifiable phenotype, such as antibiotic resistance or a complementing nutrition auxotrophic factor, which can be identified or assayed when the EMF trap vector is placed within an appropriate host under appropriate conditions. As described above, a EMF will modulate the expression of an operably linked marker sequence. A more detailed discussion of various marker sequences is provided below. A sequence which is suspected as being an EMF is cloned in all three reading frames in one or more restriction sites upstream from the marker sequence in the EMF trap vector. The vector is then transformed into an appropriate host using known procedures and the phenotype of the transformed host in examined under appropriate conditions. As described above, an EMF will modulate the expression of an operably linked marker sequence.

As used herein, a "diagnostic fragment," DF, means a series of nucleotide molecules which selectively hybridize to *Borrelia burgdorferi* sequences. DFs can be readily identified by identifying unique sequences within contigs of the *Borrelia burgdorferi* genome, such as by using well-known computer analysis software, and by generating and testing probes or

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amplification primers consisting of the DF sequence in an appropriate diagnostic format which determines amplification or hybridization selectivity.

The sequences falling within the scope of the present invention are not limited to the specific sequences herein described, but also include allelic and species variations thereof. Allelic and species variations can be routinely determined by comparing the sequences provided in SEQ ID NOS:1-155, ORF IDs and ORFs within, a representative fragment thereof, or a nucleotide sequence at least 99% and preferably 99.9% identical to SEQ ID NOS: 1-155, ORF IDs and ORFs within, with a sequence from another isolate of the same species. Furthermore, to accommodate codon variability, the invention includes nucleic acid molecules coding for the same amino acid sequences as do the specific ORFs disclosed herein. In other words, in the coding region of an ORF, substitution of one codon for another which encodes the same amino acid is expressly contemplated.

Any specific sequence disclosed herein can be readily screened for errors by resequencing a particular fragment, such as an ORF, in both directions (*i.e.*, sequence both strands). Alternatively, error screening can be performed by sequencing corresponding polynucleotides of *Borrelia burgdorferi* origin isolated by using part or all of the fragments in question as a probe or primer.

Each of the ORF IDs and ORFs of the *Borrelia burgdorferi* genome disclosed in Tables 1-6, and the EMFs found 5' to the ORF IDs, can be used as polynucleotide reagents in numerous ways. For example, the sequences can be used as diagnostic probes or diagnostic amplification primers to detect the presence of a specific microbe in a sample, particularly *Borrelia burgdorferi*. Especially preferred in this regard are ORF IDs and ORFs such as those of Tables 3 and 6, which do not match previously characterized sequences from other organisms and thus are most likely to be highly selective for *Borrelia burgdorferi*. Also particularly preferred are ORF IDs and ORFs that can be used to distinguish between strains of *Borrelia burgdorferi*, particularly those that distinguish medically important strain, such as drug-resistant strains.

In addition, the fragments of the present invention, as broadly described, can be used to control gene expression through triple helix formation or antisense DNA or RNA, both of which methods are based on the binding of a polynucleotide sequence to DNA or RNA. Triple helixformation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Information from the sequences of the present invention can be used to design antisense and triple helix-forming oligonucleotides. Polynucleotides suitable for use in these methods are usually 20 to 40 bases in length and are designed to be complementary to a region of the gene involved in transcription, for triple-helix formation, or to the mRNA itself, for antisense inhibition. Both techniques have been demonstrated to be effective in model systems, and the requisite techniques are well known and involve routine procedures. Triple helix techniques are discussed in, for example, Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991). Antisense techniques in general are discussed in, for instance, Okano,

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J. Neurochem. 56:560 (1991) and Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)).

The present invention further provides recombinant constructs comprising one or more fragments of the *Borrelia burgdorferi* genomic fragments and contigs of the present invention. Certain preferred recombinant constructs of the present invention comprise a vector, such as a plasmid or viral vector, into which a fragment of the *Borrelia burgdorferi* genome has been inserted, in a forward or reverse orientation. In the case of a vector comprising one of the ORF IDs or ORFs of the present invention, the vector may further comprise regulatory sequences, including for example, a promoter, operably linked to the ORF ID or ORF. For vectors comprising the EMFs of the present invention, the vector may further comprise a marker sequence or heterologous ORF ID or ORF operably linked to the EMF.

Large numbers of suitable vectors and promoters are known to those of skill in the art and are commercially available for generating the recombinant constructs of the present invention. The following vectors are provided by way of example. Useful bacterial vectors include phagescript, PsiX174, pBluescript SK, pBS KS, pNH8a, pNH16a, pNH18a, pNH46a (available from Stratagene); pTrc99A, pKK223-3, pKK233-3, pDR540, pRIT5 (available from Pharmacia); pQE vectors (available from Promega). Useful eukaryotic vectors include pWLneo, pSV2cat, pOG44, pXT1, pSG (available from Stratagene) pSVK3, pBPV, pMSG, pSVL (available from Pharmacia).

Promoter regions can be selected from any desired gene using CAT (chloramphenicol transferase) vectors or other vectors with selectable markers. Two appropriate vectors are pKK232-8 and pCM7. Particular named bacterial promoters include lacI, lacZ, T3, T7, gpt, lambda PR, and trc. Eukaryotic promoters include CMV immediate early, HSV thymidine kinase, early and late SV40, LTRs from retrovirus, and mouse metallothionein- I. Selection of the appropriate vector and promoter is well within the level of ordinary skill in the art.

The present invention further provides host cells containing any one of the isolated fragments of the *Borrelia burgdorferi* genomic fragments and contigs of the present invention, wherein the fragment has been introduced into the host cell using known methods. The host cell can be a higher eukaryotic host cell, such as a mammalian cell, a lower eukaryotic host cell, such as a yeast cell, or a procaryotic cell, such as a bacterial cell.

A polynucleotide of the present invention, such as a recombinant construct comprising an ORF of the present invention, may be introduced into the host by a variety of well established techniques that are standard in the art, such as calcium phosphate transfection, DEAE, dextran mediated transfection and electroporation, which are described in, for instance, Davis, L. et al., BASIC METHODS IN MOLECULAR BIOLOGY (1986).

A host cell containing one of the fragments of the *Borrelia burgdorferi* genomic fragments and contigs of the present invention, can be used in conventional manners to produce the gene product encoded by the isolated fragment (in the case of an ORF) or can be used to produce a heterologous protein under the control of the EMF.

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The present invention further provides isolated polypeptides encoded by the nucleic acid fragments of the present invention or by degenerate variants of the nucleic acid fragments of the present invention. By "degenerate variant" is intended nucleotide fragments which differ from a nucleic acid fragment of the present invention (e.g., an ORF) by nucleotide sequence but, due to the degeneracy of the Genetic Code, encode an identical polypeptide sequence.

Preferred nucleic acid fragments of the present invention are the ORF IDs depicted in Tables 2, 3, 5 and 6, and ORFs witin, which encode proteins.

A variety of methodologies known in the art can be utilized to obtain any one of the isolated polypeptides or proteins of the present invention. At the simplest level, the amino acid sequence can be synthesized using commercially available peptide synthesizers. This is particularly useful in producing small peptides and fragments of larger polypeptides. Such short fragments as may be obtained most readily by synthesis are useful, for example, in generating antibodies against the native polypeptide, as discussed further below.

In an alternative method, the polypeptide or protein is purified from bacterial cells which naturally produce the polypeptide or protein. One skilled in the art can readily employ well-known methods for isolating polypeptides and proteins to isolate and purify polypeptides or proteins of the present invention produced naturally by a bacterial strain, or by other methods. Methods for isolation and purification that can be employed in this regard include, but are not limited to, immunochromatography, HPLC, size-exclusion chromatography, ion-exchange chromatography, and immuno-affinity chromatography.

The polypeptides and proteins of the present invention also can be purified from cells which have been altered to express the desired polypeptide or protein. As used herein, a cell is said to be altered to express a desired polypeptide or protein when the cell, through genetic manipulation, is made to produce a polypeptide or protein which it normally does not produce or which the cell normally produces at a lower level. Those skilled in the art can readily adapt procedures for introducing and expressing either recombinant or synthetic sequences into eukaryotic or prokaryotic cells in order to generate a cell which produces one of the polypeptides or proteins of the present invention.

The polypeptides of the present invention are preferably provided in an isolated form, and preferably are substantially purified. A recombinantly produced version of the *B. burgdorferi* polypeptide can be substantially purified by the one-step method described by Smith et al. (1988) Gene 67:31-40. Polypeptides of the invention also can be purified from natural or recombinant sources using antibodies directed against the polypeptides of the invention in methods which are well known in the art of protein purification.

The invention further provides for isolated *B. burgdorferi* polypeptides comprising an amino acid sequence selected from the group including: (a) the amino acid sequence of a full-length *B. burgdorferi* polypeptide having the complete amino acid sequence from the first methionine codon to the termination codon of each sequence listed in SEQ ID NOS:1-155, wherein said termination codon is at the end of each SEQ ID NO: and said first methionine is the

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first methionine in frame with said termination codon; and (b) the amino acid sequence of a full-length *B. burgdorferi* polypeptide having the complete amino acid sequence in (a) excepting the N-terminal methionine.

The polypeptides of the present invention also include polypeptides having an amino acid sequence at least 80% identical, more preferably at least 90% identical, and still more preferably 95%, 96%, 97%, 98% or 99% identical to those described in (a) and (b) above.

The present invention is further directed to polynucleotides encoding portions or fragments of the amino acid sequences described herein as well as to portions or fragments of the isolated amino acid sequences described herein. Fragments include portions of the amino acid sequences described herein at least 5 contiguous amino acid in length and selected from any two integers, one of which representing an N-terminal position and another representing a C-terminal position. The initiation codon of the ORFs of the present invention is position 1. The initiation codon (positon 1) for purposes of the present invention is the first methionine codon of each ORF ID which is in frame with the termination codon at the end of each said sequence. Every combination of a N-terminal and C-terminal position that a fragment at least 5 contiguous amino acid residues in length could occupy, on any given ORF is included in the invention, i.e., from initiation codon up to the termination codon. "At least" means a fragment may be 5 contiguous amino acid residues in length or any integer between 5 and the number of residues in an ORF, minus 1. Therefore, included in the invention are contiguous fragments specified by any Nterminal and C-terminal positions of amino acid sequence set forth in SEQ ID NOS:1-155 or Tables 1-6 wherein the contiguous fragment is any integer between 5 and the number of residues in an ORF minus 1.

Further, the invention includes polypeptides comprising fragments specified by size, in amino acid residues, rather than by N-terminal and C-terminal positions. The invention includes any fragment size, in contiguous amino acid residues, selected from integers between 5 and the number of residues in an ORF, minus 1. Preferred sizes of contiguous polypeptide fragments include about 5 amino acid residues, about 10 amino acid residues, about 20 amino acid residues, about 30 amino acid residues, about 40 amino acid residues, about 50 amino acid residues, about 100 amino acid residues, about 200 amino acid residues, about 300 amino acid residues, and about 400 amino acid residues. The preferred sizes are, of course, meant to exemplify, not limit, the present invention as all size fragments representing any integer between 5 and the number of residues in a full length sequence minus 1 are included in the invention. The present invention also provides for the exclusion of any fragments specified by N-terminal and C-terminal positions or by size in amino acid residues as described above. Any number of fragments specified by N-terminal and C-terminal positions or by size in amino acid residues as described above may be excluded.

The above fragments need not be active since they would be useful, for example, in immunoassays, in epitope mapping, epitope tagging, to generate antibodies to a particular portion of the protein, as vaccines, and as molecular weight markers.

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Further polypeptides of the present invention include polypeptides which have at least 90% similarity, more preferably at least 95% similarity, and still more preferably at least 96%, 97%, 98% or 99% similarity to those described above.

A further embodiment of the invention relates to a polypeptide which comprises the amino acid sequence of a *B. burgdorferi* polypeptide having an amino acid sequence which contains at least one conservative amino acid substitution, but not more than 50 conservative amino acid substitutions, not more than 40 conservative amino acid substitutions, not more than 30 conservative amino acid substitutions, and not more than 20 conservative amino acid substitutions. Also provided are polypeptides which comprise the amino acid sequence of a *B. burgdorferi* polypeptide, having at least one, but not more than 10, 9, 8, 7, 6, 5, 4, 3, 2 or 1 conservative amino acid substitutions.

By a polypeptide having an amino acid sequence at least, for example, 95% "identical" to a query amino acid sequence of the present invention, it is intended that the amino acid sequence of the subject polypeptide is identical to the query sequence except that the subject polypeptide sequence may include up to five amino acid alterations per each 100 amino acids of the query amino acid sequence. In other words, to obtain a polypeptide having an amino acid sequence at least 95% identical to a query amino acid sequence, up to 5% of the amino acid residues in the subject sequence may be inserted, deleted, (indels) or substituted with another amino acid. These alterations of the reference sequence may occur at the amino or carboxy terminal positions of the reference amino acid sequence or anywhere between those terminal positions, interspersed either individually among residues in the reference sequence or in one or more contiguous groups within the reference sequence.

As a practical matter, whether any particular polypeptide is at least 90%, 95%, 96%, 97%, 98% or 99% identical to the ORF amino acid sequences encoded by the sequences of SEQ ID NOS:1-155, as described hererin, can be determined conventionally using known computer programs. A preferred method for determining the best overall match between a query sequence (a sequence of the present invention) and a subject sequence, also referred to as a global sequence alignment, can be determined using the FASTDB computer program based on the algorithm of Brutlag et al., (1990) Comp. App. Biosci. 6:237-245. In a sequence alignment the query and subject sequences are both amino acid sequences. The result of said global sequence alignment is in percent identity. Preferred parameters used in a FASTDB amino acid alignment are:

Matrix=PAM 0, k-tuple=2, Mismatch Penalty=1, Joining Penalty=20, Randomization Group Length=0, Cutoff Score=1, Window Size=sequence length, Gap Penalty=5, Gap Size Penalty=0.05, Window Size=500 or the length of the subject amino acid sequence, whichever is shorter.

If the subject sequence is shorter than the query sequence due to N- or C-terminal deletions, not because of internal deletions, the results, in percent identity, must be manually corrected. This is because the FASTDB program does not account for N- and C-terminal truncations of the subject sequence when calculating global percent identity. For subject

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sequences truncated at the N- and C-termini, relative to the query sequence, the percent identity is corrected by calculating the number of residues of the query sequence that are N- and C-terminal of the subject sequence, which are not matched/aligned with a corresponding subject residue, as a percent of the total bases of the query sequence. Whether a residue is matched/aligned is determined by results of the FASTDB sequence alignment. This percentage is then subtracted from the percent identity, calculated by the above FASTDB program using the specified parameters, to arrive at a final percent identity score. This final percent identity score is what is used for the purposes of the present invention. Only residues to the N- and C-termini of the subject sequence, which are not matched/aligned with the query sequence, are considered for the purposes of manually adjusting the percent identity score. That is, only query amino acid residues outside the farthest N- and C-terminal residues of the subject sequence.

For example, a 90 amino acid residue subject sequence is aligned with a 100 residue query sequence to determine percent identity. The deletion occurs at the N-terminus of the subject sequence and therefore, the FASTDB alignment does not match/align with the first 10 residues at the N-terminus. The 10 unpaired residues represent 10% of the sequence (number of residues at the N- and C- termini not matched/total number of residues in the query sequence) so 10% is subtracted from the percent identity score calculated by the FASTDB program. If the remaining 90 residues were perfectly matched the final percent identity would be 90%. In another example, a 90 residue subject sequence is compared with a 100 residue query sequence. This time the deletions are internal so there are no residues at the N- or C-termini of the subject sequence which are not matched/aligned with the query. In this case the percent identity calculated by FASTDB is not manually corrected. Once again, only residue positions outside the N- and C-terminal ends of the subject sequence, as displayed in the FASTDB alignment, which are not matched/aligned with the query sequence are manually corrected. No other manual corrections are to made for the purposes of the present invention.

The above polypeptide sequences are included irrespective of whether they have their

normal biological activity. This is because even where a particular polypeptide molecule does not have biological activity, one of skill in the art would still know how to use the polypeptide, for instance, as a vaccine or to generate antibodies. Other uses of the polypeptides of the present invention that do not have *B. burgdorferi* activity include, *inter alia*, as epitope tags, in epitope mapping, and as molecular weight markers on SDS-PAGE gels or on molecular sieve gel filtration columns using methods known to those of skill in the art.

As described below, the polypeptides of the present invention can also be used to raise polyclonal and monoclonal antibodies, which are useful in assays for detecting *B. burgdorferi* protein expression or as agonists and antagonists capable of enhancing or inhibiting *B. burgdorferi* protein function. Further, such polypeptides can be used in the yeast two-hybrid system to "capture" *B. burgdorferi* protein binding proteins which are also candidate agonists and antagonists according to the present invention. *See, e.g.*, Fields et al. (1989) Nature 340:245-246.

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Any host/vector system can be used to express one or more of the ORFs of the present invention. These include, but are not limited to, eukaryotic hosts such as HeLa cells, CV-1 cell, COS cells, and Sf9 cells, as well as prokaryotic host such as *E. coli* and *B. subtilis*. The most preferred cells are those which do not normally express the particular polypeptide or protein or which expresses the polypeptide or protein at low natural level.

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"Recombinant," as used herein, means that a polypeptide or protein is derived from recombinant (e.g., microbial or mammalian) expression systems. "Microbial" refers to recombinant polypeptides or proteins made in bacterial or fungal (e.g., yeast) expression systems. As a product, "recombinant microbial"defines a polypeptide or protein essentially free of native endogenous substances and unaccompanied by associated native glycosylation. Polypeptides or proteins expressed in most bacterial cultures, e.g., E. coli, will be free of glycosylation modifications; polypeptides or proteins expressed in yeast will have a glycosylation pattern different from that expressed in mammalian cells.

"Nucleotide sequence" refers to a heteropolymer of deoxyribonucleotides. Generally, DNA segments encoding the polypeptides and proteins provided by this invention are assembled from fragments of the *Borrelia burgdorferi* genome and short oligonucleotide linkers, or from a series of oligonucleotides, to provide a synthetic gene which is capable of being expressed in a recombinant transcriptional unit comprising regulatory elements derived from a microbial or viral operon.

Recombinant expression vehicle or vector" refers to a plasmid or phage or virus or vector, for expressing a polypeptide from a DNA (RNA) sequence. The expression vehicle can comprise a transcriptional unit comprising an assembly of (1) a genetic regulatory elements necessary for gene expression in the host, including elements required to initiate and maintain transcription at a level sufficient for suitable expression of the desired polypeptide, including, for example, promoters and, where necessary, an enhancer and a polyadenylation signal; (2) a structural or coding sequence which is transcribed into mRNA and translated into protein, and (3) appropriate signals to initiate translation at the beginning of the desired coding region and terminate translation at its end. Structural units intended for use in yeast or eukaryotic expression systems preferably include a leader sequence enabling extracellular secretion of translated protein by a host cell. Alternatively, where recombinant protein is expressed without a leader or transport sequence, it may include an N-terminal methionine residue. This residue may or may not be subsequently cleaved from the expressed recombinant protein to provide a final product.

"Recombinant expression system" means host cells which have stably integrated a recombinant transcriptional unit into chromosomal DNA or carry the recombinant transcriptional unit extra chromosomally. The cells can be prokaryotic or eukaryotic. Recombinant expression systems as defined herein will express heterologous polypeptides or proteins upon induction of the regulatory elements linked to the DNA segment or synthetic gene to be expressed.

Mature proteins can be expressed in mammalian cells, yeast, bacteria, or other cells under the control of appropriate promoters. Cell-free translation systems can also be employed to

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produce such proteins using RNAs derived from the DNA constructs of the present invention. Appropriate cloning and expression vectors for use with prokaryotic and eukaryotic hosts are described in Sambrook et al., Molecular Cloning: A Laboratory Manual, 2nd Edition, Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York (1989), the disclosure of which is hereby incorporated by reference in its entirety.

Generally, recombinant expression vectors will include origins of replication and selectable markers permitting transformation of the host cell, e.g., the ampicillin resistance gene of E. coli and S. cerevisiae TRP1 gene, and a promoter derived from a highly expressed gene to direct transcription of a downstream structural sequence. Such promoters can be derived from operons encoding glycolytic enzymes such as 3- phosphoglycerate kinase (PGK), alpha-factor, acid phosphatase, or heat shock proteins, among others. The heterologous structural sequence is assembled in appropriate phase with translation initiation and termination sequences, and preferably, a leader sequence capable of directing secretion of translated protein into the periplasmic space or extracellular medium. Optionally, the heterologous sequence can encode a fusion protein including an N-terminal identification peptide imparting desired characteristics, e.g., stabilization or simplified purification of expressed recombinant product.

Useful expression vectors for bacterial use are constructed by inserting a structural DNA sequence encoding a desired protein together with suitable translation initiation and termination signals in operable reading phase with a functional promoter. The vector will comprise one or more phenotypic selectable markers and an origin of replication to ensure maintenance of the vector and, when desirable, provide amplification within the host.

Suitable prokaryotic hosts for transformation include strains of *E. coli*, *B. subtilis*, *Salmonella typhimurium* and various species within the genera *Pseudomonas* and *Streptomyces*. Others may, also be employed as a matter of choice.

As a representative but non-limiting example, useful expression vectors for bacterial use can comprise a selectable marker and bacterial origin of replication derived from commercially available plasmids comprising genetic elements of the well known cloning vector pBR322 (ATCC 37017). Such commercial vectors include, for example, pKK223-3 (available form Pharmacia Fine Chemicals, Uppsala, Sweden) and GEM 1 (available from Promega Biotec, Madison, WI, USA). These pBR322 "backbone" sections are combined with an appropriate promoter and the structural sequence to be expressed.

Following transformation of a suitable host strain and growth of the host strain to an appropriate cell density, the selected promoter, where it is inducible, is derepressed or induced by appropriate means (e.g., temperature shift or chemical induction) and cells are cultured for an additional period to provide for expression of the induced gene product. Thereafter cells are typically harvested, generally by centrifugation, disrupted to release expressed protein, generally by physical or chemical means, and the resulting crude extract is retained for further purification.

Various mammalian cell culture systems can also be employed to express recombinant protein. Examples of mammalian expression systems include the COS-7 lines of monkey kidney

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fibroblasts, described in Gluzman, Cell 23:175 (1981), and other cell lines capable of expressing a compatible vector, for example, the C127, 3T3, CHO, HeLa and BHK cell lines.

Mammalian expression vectors will comprise an origin of replication, a suitable promoter and enhancer, and also any necessary ribosome binding sites, polyadenylation site, splice donor and acceptor sites, transcriptional termination sequences, and 5' flanking nontranscribed sequences. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early promoter, enhancer, splice, and polyadenylation sites may be used to provide the required nontranscribed genetic elements.

Recombinant polypeptides and proteins produced in bacterial culture is usually isolated by initial extraction from cell pellets, followed by one or more salting-out, aqueous ion exchange or size exclusion chromatography steps. Microbial cells employed in expression of proteins can be disrupted by any convenient method, including freeze-thaw cycling, sonication, mechanical disruption, or use of cell lysing agents. Protein refolding steps can be used, as necessary, in completing configuration of the mature protein. Finally, high performance liquid chromatography (HPLC) can be employed for final purification steps.

The present invention further includes isolated polypeptides, proteins and nucleic acid molecules which are substantially equivalent to those herein described. As used herein, substantially equivalent can refer both to nucleic acid and amino acid sequences, for example a mutant sequence, that varies from a reference sequence by one or more substitutions, deletions, or additions, the net effect of which does not result in an adverse functional dissimilarity between reference and subject sequences. Particularly preferred in this regard are conservative substitutions, known to those of skill in the art. For purposes of the present invention, sequences having equivalent biological activity, and equivalent expression characteristics are considered substantially equivalent. For purposes of determining equivalence, truncation of the mature sequence (e.g., removal of leader sequence(s)) should be disregarded.

The invention further provides methods of obtaining homologs from other strains of Borrelia burgdorferi, of the fragments of the Borrelia burgdorferi genome of the present invention and homologs of the proteins encoded by the ORFs of the present invention. As used herein, a sequence or protein of Borrelia burgdorferi is defined as a homolog of a fragment of the Borrelia burgdorferi fragments or contigs or a protein encoded by one of the ORFs of the present invention, if it shares significant homology to one of the fragments of the Borrelia burgdorferi genome of the present invention or a protein encoded by one of the ORFs of the present invention. Specifically, by using the sequence disclosed herein as a probe or as primers, and techniques such as PCR cloning and colony/plaque hybridization, one skilled in the art can obtain homologs.

As used herein, two nucleic acid molecules or proteins are said to "share significant homology" if the two contain regions which possess greater than 85% sequence (amino acid or nucleic acid) homology. Preferred homologs in this regard are those with more than 90% homology. Especially preferred are those with 95% or more homology. Among especially

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preferred homologs those with 96, 97%, 98%, 99% or more homology are particularly preferred. The most preferred homologs among these are those with 99.9% homology or more. It will be understood that, among measures of homology, identity is particularly preferred in this regard.

Region specific primers or probes derived from the nucleotide sequence provided in SEQ ID NOS: 1-155 or from a nucleotide sequence at least 95%, particularly at least 96%, 97%, 98% or 99%, especially at least 99.5% identical to a sequence of SEQ ID NOS: 1-155 can be used to prime DNA synthesis and PCR amplification, as well as to identify colonies containing cloned DNA encoding a homolog. Methods suitable to this aspect of the present invention are well known and have been described in great detail in many publications such as, for example, Innis et al., PCR Protocols, Academic Press, San Diego, CA (1990)).

When using primers derived from SEQ ID NOS: 1-155 or from a nucleotide sequence having an aforementioned identity to a sequence of SEQ ID NOS:1-155, one skilled in the art will recognize that by employing high stringency conditions (e.g., annealing at 50-60°C in 6X SSPC and 50% formamide, and washing at 50-65°C in 0.5X SSPC) only sequences which are greater than 75% homologous to the primer will be amplified. By employing lower stringency conditions (e.g., hybridizing at 35-37°C in 5X SSPC and 40-45% formamide, and washing at 42°C in 0.5X SSPC), sequences which are greater than 40-50% homologous to the primer will also be amplified.

When using DNA probes derived from SEQ ID NOS:1-155, or from a nucleotide sequence having an aforementioned identity to a sequence of SEQ ID NOS: 1-155, for colony/plaque hybridization, one skilled in the art will recognize that by employing high stringency conditions (e.g., hybridizing at 50-65°C in 5X SSPC and 50% formamide, and washing at 50-65°C in 0.5X SSPC), sequences having regions which are greater than 90% homologous to the probe can be obtained, and that by employing lower stringency conditions (e.g., hybridizing at 35-37°C in 5X SSPC and 40-45% formamide, and washing at 42°C in 0.5X SSPC), sequences having regions which are greater than 35-45% homologous to the probe will be obtained.

Any organism can be used as the source for homologs of the present invention so long as the organism naturally expresses such a protein or contains genes encoding the same. The most preferred organism for isolating homologs are bacteria which are closely related to *Borrelia burgdorferi*.

ILLUSTRATIVE USES OF COMPOSITIONS OF THE INVENTION

Each ORF of the ORF IDs provided in Tables 1, 2, 4 and 5 is identified with a function by homology to a known gene or polypeptide. As a result, one skilled in the art can use the polypeptides of the present invention for commercial, therapeutic and industrial purposes consistent with the type of putative identification of the polypeptide. Such identifications permit one skilled in the art to use the *Borrelia burgdorferi* ORFs in a manner similar to the known type

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of sequences for which the identification is made; for example, to ferment a particular sugar source or to produce a particular metabolite. A variety of reviews illustrative of this aspect of the invention are available, including the following reviews on the industrial use of enzymes, for example, BIOCHEMICAL ENGINEERING AND BIOTECHNOLOGY HANDBOOK, 2nd Ed., MacMillan Publications, Ltd. NY (1991) and BIOCATALYSTS IN ORGANIC SYNTHESES, Tramper *et al.*, Eds., Elsevier Science Publishers, Amsterdam, The Netherlands (1985). A variety of exemplary uses that illustrate this and similar aspects of the present invention are discussed below.

1. Biosynthetic Enzymes

Open reading frames encoding proteins involved in mediating the catalytic reactions involved in intermediary and macromolecular metabolism, the biosynthesis of small molecules, cellular processes and other functions includes enzymes involved in the degradation of the intermediary products of metabolism, enzymes involved in central intermediary metabolism, enzymes involved in respiration, both aerobic and anaerobic, enzymes involved in fermentation, enzymes involved in ATP proton motor force conversion, enzymes involved in broad regulatory function, enzymes involved in amino acid synthesis, enzymes involved in nucleotide synthesis, enzymes involved in cofactor and vitamin synthesis, can be used for industrial biosynthesis.

The various metabolic pathways present in *Borrelia burgdorferi* can be identified based on absolute nutritional requirements as well as by examining the various enzymes identified in Table 1-6 and SEQ ID NOS:1-155.

Of particular interest are polypeptides involved in the degradation of intermediary metabolites as well as non-macromolecular metabolism. Such enzymes include amylases, glucose oxidases, and catalase.

Proteolytic enzymes are another class of commercially important enzymes. Proteolytic enzymes find use in a number of industrial processes including the processing of flax and other vegetable fibers, in the extraction, clarification and depectinization of fruit juices, in the extraction of vegetables' oil and in the maceration of fruits and vegetables to give unicellular fruits. A detailed review of the proteolytic enzymes used in the food industry is provided in Rombouts et al., Symbiosis 21:79 (1986) and Voragen et al. in Biocatalysts In Agricultural Biotechnology, Whitaker et al., Eds., American Chemical Society Symposium Series 389:93 (1989).

The metabolism of sugars is an important aspect of the primary metabolism of *Borrelia burgdorferi*. Enzymes involved in the degradation of sugars, such as, particularly, glucose, galactose, fructose and xylose, can be used in industrial fermentation. Some of the important sugar transforming enzymes, from a commercial viewpoint, include sugar isomerases such as glucose isomerase. Other metabolic enzymes have found commercial use such as glucose oxidases which produces ketogulonic acid (KGA). KGA is an intermediate in the commercial production of ascorbic acid using the Reichstein's procedure, as described in Krueger *et al.*, *Biotechnology* <u>6(A)</u>, Rhine *et al.*, Eds., Verlag Press, Weinheim, Germany (1984).

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Glucose oxidase (GOD) is commercially available and has been used in purified form as well as in an immobilized form for the deoxygenation of beer. See, for instance, Hartmeir et al., Biotechnology Letters 1:21 (1979). The most important application of GOD is the industrial scale fermentation of gluconic acid. Market for gluconic acids which are used in the detergent, textile, leather, photographic, pharmaceutical, food, feed and concrete industry, as described, for example, in Bigelis et al., beginning on page 357 in GENE MANIPULATIONS AND FUNGI; Benett et al., Eds., Academic Press, New York (1985). In addition to industrial applications, GOD has found applications in medicine for quantitative determination of glucose in body fluids recently in biotechnology for analyzing syrups from starch and cellulose hydrosylates. This application is described in Owusu et al., Biochem. et Biophysica. Acta. 872:83 (1986), for instance.

The main sweetener used in the world today is sugar which comes from sugar beets and sugar cane. In the field of industrial enzymes, the glucose isomerase process shows the largest expansion in the market today. Initially, soluble enzymes were used and later immobilized enzymes were developed (Krueger et al., Biotechnology, The Textbook of Industrial Microbiology, Sinauer Associated Incorporated, Sunderland, Massachusetts (1990)). Today, the use of glucose- produced high fructose syrups is by far the largest industrial business using immobilized enzymes. A review of the industrial use of these enzymes is provided by Jorgensen, Starch 40:307 (1988).

Proteinases, such as alkaline serine proteinases, are used as detergent additives and thus represent one of the largest volumes of microbial enzymes used in the industrial sector. Because of their industrial importance, there is a large body of published and unpublished information regarding the use of these enzymes in industrial processes. (See Faultman *et al.*, Acid Proteases Structure Function and Biology, Tang, J., ed., Plenum Press, New York (1977) and Godfrey *et al.*, Industrial Enzymes, MacMillan Publishers, Surrey, UK (1983) and Hepner *et al.*, Report Industrial Enzymes by 1990, Hel Hepner & Associates, London (1986)).

Another class of commercially usable proteins of the present invention are the microbial lipases, described by, for instance, Macrae et al., Philosophical Transactions of the Chiral Society of London 310:227 (1985) and Poserke, Journal of the American Oil Chemist Society 61:1758 (1984). A major use of lipases is in the fat and oil industry for the production of neutral glycerides using lipase catalyzed inter-esterification of readily available triglycerides. Application of lipases include the use as a detergent additive to facilitate the removal of fats from fabrics in the course of the washing procedures.

The use of enzymes, and in particular microbial enzymes, as catalyst for key steps in the synthesis of complex organic molecules is gaining popularity at a great rate. One area of great interest is the preparation of chiral intermediates. Preparation of chiral intermediates is of interest to a wide range of synthetic chemists particularly those scientists involved with the preparation of new pharmaceuticals, agrochemicals, fragrances and flavors. (See Davies et al., Recent Advances in the Generation of Chiral Intermediates Using Enzymes, CRC Press, Boca Raton,

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Florida (1990)). The following reactions catalyzed by enzymes are of interest to organic chemists: hydrolysis of carboxylic acid esters, phosphate esters, amides and nitriles, esterification reactions, trans-esterification reactions, synthesis of amides, reduction of alkanones and oxoalkanates, oxidation of alcohols to carbonyl compounds, oxidation of sulfides to sulfoxides, and carbon bond forming reactions such as the aldol reaction.

When considering the use of an enzyme encoded by one of the ORFs of the present invention for biotransformation and organic synthesis it is sometimes necessary to consider the respective advantages and disadvantages of using a microorganism as opposed to an isolated enzyme. Pros and cons of using a whole cell system on the one hand or an isolated partially purified enzyme on the other hand, has been described in detail by Bud *et al.*, Chemistry in Britain (1987), p. 127.

Amino transferases, enzymes involved in the biosynthesis and metabolism of amino acids, are useful in the catalytic production of amino acids. The advantages of using microbial based enzyme systems is that the amino transferase enzymes catalyze the stereo- selective synthesis of only L-amino acids and generally possess uniformly high catalytic rates. A description of the use of amino transferases for amino acid production is provided by Roselle-David, *Methods of Enzymology 136*:479 (1987).

Another category of useful proteins encoded by the ORFs of the present invention include enzymes involved in nucleic acid synthesis, repair, and recombination.

2. Generation of Antibodies

As described here, the proteins of the present invention, as well as homologs thereof, can be used in a variety of procedures and methods known in the art which are currently applied to other proteins. The proteins of the present invention can further be used to generate an antibody which selectively binds the protein.

B. burgdorferi protein-specific antibodies for use in the present invention can be raised against the intact B. burgdorferi protein or an antigenic polypeptide fragment thereof, which may be presented together with a carrier protein, such as an albumin, to an animal system (such as rabbit or mouse) or, if it is long enough (at least about 25 amino acids), without a carrier.

As used herein, the term "antibody" (Ab) or "monoclonal antibody" (Mab) is meant to include intact molecules, single chain whole antibodies, and antibody fragments. Antibody fragments of the present invention include Fab and F(ab')2 and other fragments including single-chain Fvs (scFv) and disulfide-linked Fvs (sdFv). Also included in the present invention are chimeric and humanized monoclonal antibodies and polyclonal antibodies specific for the polypeptides of the present invention. The antibodies of the present invention may be prepared by any of a variety of methods. For example, cells expressing a polypeptide of the present invention or an antigenic fragment thereof can be administered to an animal in order to induce the production of sera containing polyclonal antibodies. For example, a preparation of *B. burgdorferi* polypeptide or fragment thereof is prepared and purified to render it substantially free

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of natural contaminants. Such a preparation is then introduced into an animal in order to produce polyclonal antisera of greater specific activity.

In a preferred method, the antibodies of the present invention are monoclonal antibodies or binding fragments thereof. Such monoclonal antibodies can be prepared using hybridoma technology. *See, e.g.*, Harlow et al., ANTIBODIES: A LABORATORY MANUAL, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988); Hammerling, et al., in: MONOCLONAL ANTIBODIES AND T-CELL HYBRIDOMAS 563-681 (Elsevier, N.Y., 1981). Fab and F(ab')2 fragments may be produced by proteolytic cleavage, using enzymes such as papain (to produce Fab fragments) or pepsin (to produce F(ab')2 fragments). Alternatively, *B. burgdorferi* polypeptide-binding fragments, chimeric, and humanized antibodies can be produced through the application of recombinant DNA technology or through synthetic chemistry using methods known in the art.

Alternatively, additional antibodies capable of binding to the polypeptide antigen of the present invention may be produced in a two-step procedure through the use of anti-idiotypic antibodies. Such a method makes use of the fact that antibodies are themselves antigens, and that, therefore, it is possible to obtain an antibody which binds to a second antibody. In accordance with this method, *B. burgdorferi* polypeptide-specific antibodies are used to immunize an animal, preferably a mouse. The splenocytes of such an animal are then used to produce hybridoma cells, and the hybridoma cells are screened to identify clones which produce an antibody whose ability to bind to the *B. burgdorferi* polypeptide-specific antibody can be blocked by the *B. burgdorferi* polypeptide antigen. Such antibodies comprise anti-idiotypic antibodies to the *B. burgdorferi* polypeptide-specific antibody and can be used to immunize an animal to induce formation of further *B. burgdorferi* polypeptide-specific antibodies.

Antibodies and fragements thereof of the present invention may be described by the portion of a polypeptide of the present invention recognized or specifically bound by the antibody. Antibody binding fragements of a polypeptide of the present invention may be described or specified in the same manner as for polypeptide fragements discussed above., i.e, by N-terminal and C-terminal positions or by size in contiguous amino acid residues. Any number of antibody binding fragments, of a polypeptide of the present invention, specified by N-terminal and C-terminal positions or by size in amino acid residues, as described above, may also be excluded from the present invention. Therefore, the present invention includes antibodies the specifically bind a particuarlly discribed fragement of a polypeptide of the present invention and allows for the exclusion of the same.

Antibodies and fragements thereof of the present invention may also be described or specified in terms of their cross-reactivity. Antibodies and fragements that do not bind polypeptides of any other species of *Borrelia* other than *B. burgdorferi* are included in the present invention. Likewise, antibodies and fragements that bind only species of *Borrelia*, i.e. antibodies and fragements that do not bind bacteria from any genus other than *Borrelia*, are included in the present invention.

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3. Epitope-Bearing Portions

In another aspect, the invention provides peptides and polypeptides comprising epitope-bearing portions of the *B. burgdorferi* polypeptides of the present invention. These epitopes are immunogenic or antigenic epitopes of the polypeptides of the present invention. An "immunogenic epitope" is defined as a part of a protein that elicits an antibody response when the whole protein or polypeptide is the immunogen. These immunogenic epitopes are believed to be confined to a few loci on the molecule. On the other hand, a region of a protein molecule to which an antibody can bind is defined as an "antigenic determinant" or "antigenic epitope." The number of immunogenic epitopes of a protein generally is less than the number of antigenic epitopes. *See, e.g.,* Geysen, et al. (1983) Proc. Natl. Acad. Sci. USA 81:3998- 4002. Amino acid residues comprising anigenic epitopes may be determined by algorithms such as the the Jameson-Wolf analysis or similar algorithms or by *in vivo* testing for an antigenic response using the methods described herein or those known in the art.

As to the selection of peptides or polypeptides bearing an antigenic epitope (*i.e.*, that contain a region of a protein molecule to which an antibody can bind), it is well known in that art that relatively short synthetic peptides that mimic part of a protein sequence are routinely capable of eliciting an antiserum that reacts with the partially mimicked protein. *See*, *e.g.*, Sutcliffe, et al., (1983) Science 219:660-666. Peptides capable of eliciting protein-reactive sera are frequently represented in the primary sequence of a protein, can be characterized by a set of simple chemical rules, and are confined neither to immunodominant regions of intact proteins (*i.e.*, immunogenic epitopes) nor to the amino or carboxyl terminals. Peptides that are extremely hydrophobic and those of six or fewer residues generally are ineffective at inducing antibodies that bind to the mimicked protein; longer, peptides, especially those containing proline residues, usually are effective. *See*, Sutcliffe, et al., *supra*, p. 661. For instance, 18 of 20 peptides designed according to these guidelines, containing 8-39 residues covering 75% of the sequence of the influenza virus hemagglutinin HA1 polypeptide chain, induced antibodies that reacted with the HA1 protein or intact virus; and 12/12 peptides from the MuLV polymerase and 18/18 from the rabies glycoprotein induced antibodies that precipitated the respective proteins.

Antigenic epitope-bearing peptides and polypeptides of the invention are therefore useful to raise antibodies, including monoclonal antibodies, that bind specifically to a polypeptide of the invention. Thus, a high proportion of hybridomas obtained by fusion of spleen cells from donors immunized with an antigen epitope-bearing peptide generally secrete antibody reactive with the native protein. *See* Sutcliffe, et al., *supra*, p. 663. The antibodies raised by antigenic epitope-bearing peptides or polypeptides are useful to detect the mimicked protein, and antibodies to different peptides may be used for tracking the fate of various regions of a protein precursor which undergoes post-translational processing. The peptides and anti-peptide antibodies may be used in a variety of qualitative or quantitative assays for the mimicked protein, for instance in competition assays since it has been shown that even short peptides (*e.g.*, about 9 amino acids)

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can bind and displace the larger peptides in immunoprecipitation assays. See, e.g., Wilson, et al., (1984) Cell 37:767-778. The anti-peptide antibodies of the invention also are useful for purification of the mimicked protein, for instance, by adsorption chromatography using methods known in the art.

Antigenic epitope-bearing peptides and polypeptides of the invention designed according to the above guidelines preferably contain a sequence of at least seven, more preferably at least nine and most preferably between about 10 to about 50 amino acids (i.e. any integer between 7 and 50) contained within the amino acid sequence of a polypeptide of the invention. However, peptides or polypeptides comprising a larger portion of an amino acid sequence of a polypeptide of the invention, containing about 50 to about 100 amino acids, or any length up to and including the entire amino acid sequence of a polypeptide of the invention, also are considered epitope-bearing peptides or polypeptides of the invention and also are useful for inducing antibodies that react with the mimicked protein. Preferably, the amino acid sequence of the epitope-bearing peptide is selected to provide substantial solubility in aqueous solvents (i.e., the sequence includes relatively hydrophilic residues and highly hydrophobic sequences are preferably avoided); and sequences containing proline residues are particularly preferred.

The epitope-bearing peptides and polypeptides of the present invention may be produced by any conventional means for making peptides or polypeptides including recombinant means using nucleic acid molecules of the invention. For instance, an epitope-bearing amino acid sequence of the present invention may be fused to a larger polypeptide which acts as a carrier during recombinant production and purification, as well as during immunization to produce anti-peptide antibodies. Epitope-bearing peptides also may be synthesized using known methods of chemical synthesis. For instance, Houghten has described a simple method for synthesis of large numbers of peptides, such as 10-20 mg of 248 different 13 residue peptides representing single amino acid variants of a segment of the HA1 polypeptide which were prepared and characterized (by ELISA-type binding studies) in less than four weeks (Houghten, R. A. Proc. Natl. Acad. Sci. USA 82:5131-5135 (1985)). This "Simultaneous Multiple Peptide Synthesis (SMPS)" process is further described in U.S. Patent No. 4,631,211 to Houghten and coworkers (1986). In this procedure the individual resins for the solid-phase synthesis of various peptides are contained in separate solvent-permeable packets, enabling the optimal use of the many identical repetitive steps involved in solid-phase methods. A completely manual procedure allows 500-1000 or more syntheses to be conducted simultaneously (Houghten et al. (1985) Proc. Natl. Acad. Sci. 82:5131-5135 at 5134.

Epitope-bearing peptides and polypeptides of the invention are used to induce antibodies according to methods well known in the art. See, e.g., Sutcliffe, et al., supra;; Wilson, et al., supra;; and Bittle, et al. (1985) J. Gen. Virol. 66:2347-2354. Generally, animals may be immunized with free peptide; however, anti-peptide antibody titer may be boosted by coupling of the peptide to a macromolecular carrier, such as keyhole limpet hemacyanin (KLH) or tetanus toxoid. For instance, peptides containing cysteine may be coupled to carrier using a linker such

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as m-maleimidobenzoyl-N-hydroxysuccinimide ester (MBS), while other peptides may be coupled to carrier using a more general linking agent such as glutaraldehyde. Animals such as rabbits, rats and mice are immunized with either free or carrier-coupled peptides, for instance, by intraperitoneal and/or intradermal injection of emulsions containing about 100 µg peptide or carrier protein and Freund's adjuvant. Several booster injections may be needed, for instance, at intervals of about two weeks, to provide a useful titer of anti-peptide antibody which can be detected, for example, by ELISA assay using free peptide adsorbed to a solid surface. The titer of anti-peptide antibodies in serum from an immunized animal may be increased by selection of anti-peptide antibodies, for instance, by adsorption to the peptide on a solid support and elution of the selected antibodies according to methods well known in the art.

Immunogenic epitope-bearing peptides of the invention, i.e., those parts of a protein that elicit an antibody response when the whole protein is the immunogen, are identified according to methods known in the art. For instance, Geysen, et al., supra, discloses a procedure for rapid concurrent synthesis on solid supports of hundreds of peptides of sufficient purity to react in an ELISA. Interaction of synthesized peptides with antibodies is then easily detected without removing them from the support. In this manner a peptide bearing an immunogenic epitope of a desired protein may be identified routinely by one of ordinary skill in the art. For instance, the immunologically important epitope in the coat protein of foot-and-mouth disease virus was located by Geysen et al. supra with a resolution of seven amino acids by synthesis of an overlapping set of all 208 possible hexapeptides covering the entire 213 amino acid sequence of the protein. Then, a complete replacement set of peptides in which all 20 amino acids were substituted in turn at every position within the epitope were synthesized, and the particular amino acids conferring specificity for the reaction with antibody were determined. Thus, peptide analogs of the epitope-bearing peptides of the invention can be made routinely by this method. U.S. Patent No. 4,708,781 to Geysen (1987) further describes this method of identifying a peptide bearing an immunogenic epitope of a desired protein.

Further still, U.S. Patent No. 5,194,392, to Geysen (1990), describes a general method of detecting or determining the sequence of monomers (amino acids or other compounds) which is a topological equivalent of the epitope (*i.e.*, a "mimotope") which is complementary to a particular paratope (antigen binding site) of an antibody of interest. More generally, U.S. Patent No. 4,433,092, also to Geysen (1989), describes a method of detecting or determining a sequence of monomers which is a topographical equivalent of a ligand which is complementary to the ligand binding site of a particular receptor of interest. Similarly, U.S. Patent No. 5,480,971 to Houghten, R. A. *et al.* (1996) discloses linear C₁-C₇-alkyl peralkylated oligopeptides and sets and libraries of such peptides, as well as methods for using such oligopeptide sets and libraries for determining the sequence of a peralkylated oligopeptide that preferentially binds to an acceptor molecule of interest. Thus, non-peptide analogs of the epitope-bearing peptides of the invention also can be made routinely by these methods. The entire disclosure of each document cited in this section on "Polypeptides and Fragments" is

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hereby incorporated herein by reference.

As one of skill in the art will appreciate, the polypeptides of the present invention and the epitope-bearing fragments thereof described above can be combined with parts of the constant domain of immunoglobulins (IgG), resulting in chimeric polypeptides. These fusion proteins facilitate purification and show an increased half-life *in vivo*. This has been shown, *e.g.*, for chimeric proteins consisting of the first two domains of the human CD4-polypeptide and various domains of the constant regions of the heavy or light chains of mammalian immunoglobulins. (EPA 0,394,827; Traunecker et al. (1988) Nature 331:84-86. Fusion proteins that have a disulfide-linked dimeric structure due to the IgG part can also be more efficient in binding and neutralizing other molecules than a monomeric *B. burgdorferi* polypeptide or fragment thereof alone. *See* Fountoulakis et al. (1995) J. Biochem. 270:3958-3964. Nucleic acids encoding the above epitopes of *B. burgdorferi* polypeptides can also be recombined with a gene of interest as an epitope tag to aid in detection and purification of the expressed polypeptide.

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4. Diagnostic Assays and Kits

The present invention further relates to methods for assaying Borrelia infection in an animal by detecting the expression of genes encoding Borrelia polypeptides of the present invention. The methods comprise analyzing tissue or body fluid from the animal for *Borrelia*-specific antibodies, nucleic acids, or proteins. Analysis of nucleic acid specific to *Borrelia* is assayed by PCR or hybridization techniques using nucleic acid sequences of the present invention as either hybridization probes or primers. *See, e.g.,* Sambrook et al. Molecular cloning: A Laboratory Manual (Cold Spring Harbor Laboratory Press, 2nd ed., 1989, page 54 reference); Eremeeva et al. (1994) J. Clin. Microbiol. 32:803-810 (describing differentiation among spotted fever group *Rickettsiae* species by analysis of restriction fragment length polymorphism of PCR-amplified DNA) and Chen et al. 1994 J. Clin. Microbiol. 32:589-595 (detecting *B burgdorferi* nucleic acids *via* PCR).

Where diagnosis of a disease state related to infection with *Borrelia* has already been made, the present invention is useful for monitoring progression or regression of the disease state whereby patients exhibiting enhanced *Borrelia* gene expression will experience a worse clinical outcome relative to patients expressing these gene(s) at a lower level.

By "biological sample" is intended any biological sample obtained from an animal, cell line, tissue culture, or other source which contains *Borrelia* polypeptide, mRNA, or DNA. Biological samples include body fluids (such as saliva, blood, plasma, urine, mucus, synovial fluid, etc.) tissues (such as muscle, skin, and cartilage) and any other biological source suspected of containing *Borrelia* polypeptides or nucleic acids. Methods for obtaining biological samples such as tissue are well known in the art.

The present invention is useful for detecting diseases related to *Borrelia* infections in animals. Preferred animals include monkeys, apes, cats, dogs, birds, cows, pigs, mice, horses, rabbits and humans. Particularly preferred are humans.

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Total RNA can be isolated from a biological sample using any suitable technique such as the single-step guanidinium-thiocyanate-phenol-chloroform method described in Chomczynski et al. (1987) Anal. Biochem. 162:156-159. mRNA encoding *Borrelia* polypeptides having sufficient homology to the nucleic acid sequences identified in SEQ ID NOS:1-155 to allow for hybridization between complementary sequences are then assayed using any appropriate method. These include Northern blot analysis, S1 nuclease mapping, the polymerase chain reaction (PCR), reverse transcription in combination with the polymerase chain reaction (RT-PCR), and reverse transcription in combination with the ligase chain reaction (RT-LCR).

Northern blot analysis can be performed as described in Harada et al. (1990) Cell 63:303-312. Briefly, total RNA is prepared from a biological sample as described above. For the Northern blot, the RNA is denatured in an appropriate buffer (such as glyoxal/dimethyl sulfoxide/sodium phosphate buffer), subjected to agarose gel electrophoresis, and transferred onto a nitrocellulose filter. After the RNAs have been linked to the filter by a UV linker, the filter is prehybridized in a solution containing formamide, SSC, Denhardt's solution, denatured salmon sperm, SDS, and sodium phosphate buffer. A *B. burgdorferi* polynucleotide sequence shown in SEQ ID NOS:1-155 labeled according to any appropriate method (such as the ³²P-multiprimed DNA labeling system (Amersham)) is used as probe. After hybridization overnight, the filter is washed and exposed to x-ray film. DNA for use as probe according to the present invention is described in the sections above and will preferably at least 15 nucleotides in length.

S1 mapping can be performed as described in Fujita et al. (1987) Cell 49:357-367. To prepare probe DNA for use in S1 mapping, the sense strand of an above-described *B. burgdorferi* DNA sequence of the present invention is used as a template to synthesize labeled antisense DNA. The antisense DNA can then be digested using an appropriate restriction endonuclease to generate further DNA probes of a desired length. Such antisense probes are useful for visualizing protected bands corresponding to the target mRNA (*i.e.*, mRNA encoding *Borrelia* polypeptides).

Levels of mRNA encoding *Borrelia* polypeptides are assayed, for *e.g.*, using the RT-PCR method described in Makino et al. (1990) Technique 2:295-301. By this method, the radioactivities of the "amplicons" in the polyacrylamide gel bands are linearly related to the initial concentration of the target mRNA. Briefly, this method involves adding total RNA isolated from a biological sample in a reaction mixture containing a RT primer and appropriate buffer. After incubating for primer annealing, the mixture can be supplemented with a RT buffer, dNTPs, DTT, RNase inhibitor and reverse transcriptase. After incubation to achieve reverse transcription of the RNA, the RT products are then subject to PCR using labeled primers. Alternatively, rather than labeling the primers, a labeled dNTP can be included in the PCR reaction mixture. PCR amplification can be performed in a DNA thermal cycler according to conventional techniques. After a suitable number of rounds to achieve amplification, the PCR reaction mixture is electrophoresed on a polyacrylamide gel. After drying the gel, the radioactivity of the appropriate

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bands (corresponding to the mRNA encoding the *Borrelia* polypeptides of the present invention) are quantified using an imaging analyzer. RT and PCR reaction ingredients and conditions, reagent and gel concentrations, and labeling methods are well known in the art. Variations on the RT-PCR method will be apparent to the skilled artisan. Other PCR methods that can detect the nucleic acid of the present invention can be found in PCR PRIMER: A LABORATORY MANUAL (C.W. Dieffenbach et al. eds., Cold Spring Harbor Lab Press, 1995).

The polynucleotides of the present invention, including both DNA and RNA, may be used to detect polynucleotides of the present invention or Borrelia species including B. burgdorferi using bio chip technology. The present invention includes both high density chip arrays (>1000 oligonucleotides per cm²) and low density chip arrays (<1000 oligonucleotides per cm²). Bio chips comprising arrays of polynucleotides of the present invention may be used to detect Borrelia species, including B. burgdorferi, in biological and environmental samples and to diagnose an animal, including humans, with an B. burgdorferi or other Borrelia infection. The bio chips of the present invention may comprise polynucleotide sequences of other pathogens including bacteria, viral, parasitic, and fungal polynucleotide sequences, in addition to the polynucleotide sequences of the present invention, for use in rapid diffenertial pathogenic detection and diagnosis. The bio chips can also be used to monitor an B. burgdorferi or other Borrelia infections and to monitor the genetic changes (deletions, insertions, mismatches, etc.) in response to drug therapy in the clinic and drug development in the laboratory. The bio chip technology comprising arrays of polynucleotides of the present invention may also be used to simultaneously monitor the expression of a multiplicity of genes, including those of the present invention. The polynucleotides used to comprise a selected array may be specified in the same manner as for the fragements, i.e, by their 5' and 3' positions or length in contigious base pairs and include from. Methods and particular uses of the polynucleotides of the present invention to detect Borrelia species, including B. burgdorferi, using bio chip technology include those known in the art and those of: U.S. Patent Nos. 5510270, 5545531, 5445934, 5677195, 5532128, 5556752, 5527681, 5451683, 5424186, 5607646, 5658732 and World Patent Nos. WO/9710365, WO/9511995, WO/9743447, WO/9535505, each incorporated herein in their entireties.

Biosensors using the polynucleotides of the present invention may also be used to detect, diagnose, and monitor *B. burgdorferi* or other Borrelia species and infections thereof. Biosensors using the polynucleotides of the present invention may also be used to detect particular polynucleotides of the present invention. Biosensors using the polynucleotides of the present invention may also be used to monitor the genetic changes (deletions, insertions, mismatches, etc.) in response to drug therapy in the clinic and drug development in the laboratory. Methods and particular uses of the polynucleotides of the present invention to detect Borrelia species, including *B. burgdorferi*, using biosenors include those known in the art and those of: U.S. Patent Nos 5721102, 5658732, 5631170, and World Patent Nos. WO97/35011, WO/9720203, each incorporated herein in their entireties.

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Thus, the present invention includes both bio chips and biosensors comprising polynucleotides of the present invention and methods of their use.

Assaying *Borrelia* polypeptide levels in a biological sample can occur using any art-known method, such as antibody-based techniques. For example, *Borrelia* polypeptide expression in tissues can be studied with classical immunohistological methods. In these, the specific recognition is provided by the primary antibody (polyclonal or monoclonal) but the secondary detection system can utilize fluorescent, enzyme, or other conjugated secondary antibodies. As a result, an immunohistological staining of tissue section for pathological examination is obtained. Tissues can also be extracted, *e.g.*, with urea and neutral detergent, for the liberation of *Borrelia* polypeptides for Western-blot or dot/slot assay. *See*, *e.g.*, Jalkanen, M. et al. (1985) J. Cell. Biol. 101:976-985; Jalkanen, M. et al. (1987) J. Cell . Biol. 105:3087-3096. In this technique, which is based on the use of cationic solid phases, quantitation of a *Borrelia* polypeptide can be accomplished using an isolated *Borrelia* polypeptide as a standard. This technique can also be applied to body fluids.

Other antibody-based methods useful for detecting *Borrelia* polypeptide gene expression include immunoassays, such as the ELISA and the radioimmunoassay (RIA). For example, a *Borrelia* polypeptide-specific monoclonal antibodies can be used both as an immunoabsorbent and as an enzyme-labeled probe to detect and quantify a *Borrelia* polypeptide. The amount of a *Borrelia* polypeptide present in the sample can be calculated by reference to the amount present in a standard preparation using a linear regression computer algorithm. Such an ELISA is described in Iacobelli et al. (1988) Breast Cancer Research and Treatment 11:19-30. In another ELISA assay, two distinct specific monoclonal antibodies can be used to detect *Borrelia* polypeptides in a body fluid. In this assay, one of the antibodies is used as the immunoabsorbent and the other as the enzyme-labeled probe.

The above techniques may be conducted essentially as a "one-step" or "two-step" assay. The "one-step" assay involves contacting the *Borrelia* polypeptide with immobilized antibody and, without washing, contacting the mixture with the labeled antibody. The "two-step" assay involves washing before contacting the mixture with the labeled antibody. Other conventional methods may also be employed as suitable. It is usually desirable to immobilize one component of the assay system on a support, thereby allowing other components of the system to be brought into contact with the component and readily removed from the sample. Variations of the above and other immunological methods included in the present invention can also be found in Harlow et al., ANTIBODIES: A LABORATORY MANUAL, (Cold Spring Harbor Laboratory Press, 2nd ed. 1988).

Suitable enzyme labels include, for example, those from the oxidase group, which catalyze the production of hydrogen peroxide by reacting with substrate. Glucose oxidase is particularly preferred as it has good stability and its substrate (glucose) is readily available. Activity of an oxidase label may be assayed by measuring the concentration of hydrogen peroxide formed by the enzyme-labeled antibody/substrate reaction. Besides enzymes, other suitable

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labels include radioisotopes, such as iodine (¹²⁵I, ¹²¹I), carbon (¹⁴C), sulphur (³⁵S), tritium (³H), indium (¹¹²In), and technetium (^{99m}Tc), and fluorescent labels, such as fluorescein and rhodamine, and biotin.

Further suitable labels for the *Borrelia* polypeptide-specific antibodies of the present invention are provided below. Examples of suitable enzyme labels include malate dehydrogenase, Borrelia nuclease, delta-5-steroid isomerase, yeast-alcohol dehydrogenase, alpha-glycerol phosphate dehydrogenase, triose phosphate isomerase, peroxidase, alkaline phosphatase, asparaginase, glucose oxidase, beta-galactosidase, ribonuclease, urease, catalase, glucose-6-phosphate dehydrogenase, glucoamylase, and acetylcholine esterase.

Examples of suitable radioisotopic labels include ³H, ¹¹¹In, ¹²⁵I, ¹³¹I, ³²P, ³⁵S, ¹⁴C, ⁵¹Cr, ⁵⁷To, ⁵⁸Co, ⁵⁹Fe, ⁷⁵Se, ¹⁵²Eu, ⁹⁰Y, ⁶⁷Cu, ²¹⁷Ci, ²¹¹At, ²¹²Pb, ⁴⁷Sc, ¹⁰⁹Pd, etc. ¹¹¹In is a preferred isotope where *in vivo* imaging is used since its avoids the problem of dehalogenation of the ¹²⁵I or ¹³¹I-labeled monoclonal antibody by the liver. In addition, this radionucleotide has a more favorable gamma emission energy for imaging. *See, e.g.*, Perkins et al. (1985) Eur. J. Nucl.

Med. 10:296-301; Carasquillo et al. (1987) J. Nucl. Med. 28:281-287. For example, ¹¹¹In coupled to monoclonal antibodies with 1-(P-isothiocyanatobenzyl)-DPTA has shown little uptake in non-tumors tissues, particularly the liver, and therefore enhances specificity of tumor localization. See, Esteban et al. (1987) J. Nucl. Med. 28:861-870.

Examples of suitable non-radioactive isotopic labels include ¹⁵⁷Gd, ⁵⁵Mn, ¹⁶²Dy, ⁵²Tr, and ⁵⁶Fe.

Examples of suitable fluorescent labels include an ¹⁵²Eu label, a fluorescein label, an isothiocyanate label, a rhodamine label, a phycocrythrin label, a phycocyanin label, an allophycocyanin label, an o-phthaldehyde label, and a fluorescamine label.

Examples of suitable toxin labels include, *Pseudomonas* toxin, diphtheria toxin, ricin, and cholera toxin.

Examples of chemiluminescent labels include a luminal label, an isoluminal label, an aromatic acridinium ester label, an imidazole label, an acridinium salt label, an oxalate ester label, a luciferin label, a luciferase label, and an aequorin label.

Examples of nuclear magnetic resonance contrasting agents include heavy metal nuclei such as Gd, Mn, and iron.

Typical techniques for binding the above-described labels to antibodies are provided by Kennedy et al. (1976) Clin. Chim. Acta 70:1-31, and Schurs et al. (1977) Clin. Chim. Acta 81:1-40. Coupling techniques mentioned in the latter are the glutaraldehyde method, the periodate method, the dimaleimide method, the m-maleimidobenzyl-N-hydroxy-succinimide ester method, all of which methods are incorporated by reference herein.

In a related aspect, the invention includes a diagnostic kit for use in screening serum containing antibodies specific against *B. burgdorferi* infection. Such a kit may include an isolated *B. burgdorferi* antigen comprising an epitope which is specifically immunoreactive with at least one anti-*B. burgdorferi* antibody. Such a kit also includes means for detecting the

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binding of said antibody to the antigen. In specific embodiments, the kit may include a recombinantly produced or chemically synthesized peptide or polypeptide antigen. The peptide or polypeptide antigen may be attached to a solid support.

In a more specific embodiment, the detecting means of the above-described kit includes a solid support to which said peptide or polypeptide antigen is attached. Such a kit may also include a non-attached reporter-labeled anti-human antibody. In this embodiment, binding of the antibody to the *B. burgdorferi* antigen can be detected by binding of the reporter labeled antibody to the anti-*B. burgdorferi* polypeptide antibody.

In a related aspect, the invention includes a method of detecting *B. burgdorferi* infection in a subject. This detection method includes reacting a body fluid, preferably serum, from the subject with an isolated *B. burgdorferi* antigen, and examining the antigen for the presence of bound antibody. In a specific embodiment, the method includes a polypeptide antigen attached to a solid support, and serum is reacted with the support. Subsequently, the support is reacted with a reporter-labeled anti-human antibody. The support is then examined for the presence of reporter-labeled antibody.

The solid surface reagent employed in the above assays and kits is prepared by known techniques for attaching protein material to solid support material, such as polymeric beads, dip sticks, 96-well plates or filter material. These attachment methods generally include non-specific adsorption of the protein to the support or covalent attachment of the protein, typically through a free amine group, to a chemically reactive group on the solid support, such as an activated carboxyl, hydroxyl, or aldehyde group. Alternatively, streptavidin coated plates can be used in conjunction with biotinylated antigen(s).

The polypeptides and antibodies of the present invention, including fragments thereof, may be used to detect Borrelia species including *B. burgdorferi* using bio chip and biosensor technology. Bio chip and biosensors of the present invention may comprise the polypeptides of the present invention to detect antibodies, which specifically recognize Borrelia species, including *B. burgdorferi*. Bio chip and biosensors of the present invention may also comprise antibodies which specifically recognize the polypeptides of the present invention to detect Borrelia species, including *B. burgdorferi* or specific polypeptides of the present invention. Bio chips or biosensors comprising polypeptides or antibodies of the present invention may be used to detect Borrelia species, including *B. burgdorferi*, in biological and environmental samples and to diagnose an animal, including humans, with an *B. burgdorferi* or other Borrelia infection. Thus, the present invention includes both bio chips and biosensors comprising polypeptides or antibodies of the present invention and methods of their use.

The bio chips of the present invention may further comprise polypeptide sequences of other pathogens including bacteria, viral, parasitic, and fungal polypeptide sequences, in addition to the polypeptide sequences of the present invention, for use in rapid differential pathogenic detection and diagnosis. The bio chips of the present invention may further comprise antibodies or fragements thereof specific for other pathogens including bacteria, viral, parasitic, and fungal

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polypeptide sequences, in addition to the antibodies or fragements thereof of the present invention, for use in rapid diffenertial pathogenic detection and diagnosis. The bio chips and biosensors of the present invention may also be used to monitor an B. burgdorferi or other Borrelia infection and to monitor the genetic changes (amio acid deletions, insertions, substitutions, etc.) in response to drug therapy in the clinic and drug development in the 5 laboratory. The bio chip and biosensors comprising polypeptides or antibodies of the present invention may also be used to simultaneously monitor the expression of a multiplicity of polypeptides, including those of the present invention. The polypeptides used to comprise a bio chip or biosensor of the present invention may be specified in the same manner as for the 10 fragements, i.e, by their N-terminal and C-terminal positions or length in contigious amino acid residue. Methods and particular uses of the polypeptides and antibodies of the present invention to detect Borrelia species, including B. burgdorferi, or specific polypeptides using bio chip and biosensor technology include those known in the art, those of the U.S. Patent Nos. and World Patent Nos. listed above for bio chips and biosensors using polynucleotides of the present invention, and those of: U.S. Patent Nos. 5658732, 5135852, 5567301, 5677196, 5690894 15 and World Patent Nos. WO9729366, WO9612957, each incorporated herein in their entireties.

5. Screening Assay for Binding Agents

Using the isolated proteins of the present invention, the present invention further provides methods of obtaining and identifying agents which bind to a protein encoded by one of the ORFs of the present invention or to one of the fragments and the *Borrelia burgdorferi* fragment and contigs herein described.

In general, such methods comprise steps of:

- (a) contacting an agent with an isolated protein encoded by one of the ORFs of the present invention, or an isolated fragment of the *Borrelia burgdorferi* genome; and
 - (b) determining whether the agent binds to said protein or said fragment.

The agents screened in the above assay can be, but are not limited to, peptides, carbohydrates, vitamin derivatives, or other pharmaceutical agents. The agents can be selected and screened at random or rationally selected or designed using protein modeling techniques.

For random screening, agents such as peptides, carbohydrates, pharmaceutical agents and the like are selected at random and are assayed for their ability to bind to the protein encoded by the ORF of the present invention.

Alternatively, agents may be rationally selected or designed. As used herein, an agent is said to be "rationally selected or designed" when the agent is chosen based on the configuration of the particular protein. For example, one skilled in the art can readily adapt currently available procedures to generate peptides, pharmaceutical agents and the like capable of binding to a specific peptide sequence in order to generate rationally designed antipeptide peptides, for example see Hurby *et al.*, "Application of Synthetic Peptides: Antisense Peptides," in *Synthetic*

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Peptides, A User's Guide, W. H. Freeman, NY (1992), pp. 289-307, and Kaspczak et al., Biochemistry 28:9230-8 (1989), or pharmaceutical agents, or the like.

In addition to the foregoing, one class of agents of the present invention, as broadly described, can be used to control gene expression through binding to one of the ORFs or EMFs of the present invention. As described above, such agents can be randomly screened or rationally designed/selected. Targeting the ORF or EMF allows a skilled artisan to design sequence specific or element specific agents, modulating the expression of either a single ORF or multiple ORFs which rely on the same EMF for expression control.

One class of DNA binding agents are agents which contain base residues which hybridize or form a triple helix by binding to DNA or RNA. Such agents can be based on the classic phosphodiester, ribonucleic acid backbone, or can be a variety of sulfhydryl or polymeric derivatives which have base attachment capacity.

Agents suitable for use in these methods usually contain 20 to 40 bases and are designed to be complementary to a region of the gene involved in transcription (triple helix - see Lee et al., Nucl. Acids Res. 6:3073 (1979); Cooney et al., Science 241:456 (1988); and Dervan et al., Science 251:1360 (1991)) or to the mRNA itself (antisense - Okano, J. Neurochem. 56:560 (1991); Oligodeoxynucleotides as Antisense Inhibitors of Gene Expression, CRC Press, Boca Raton, FL (1988)). Triple helix- formation optimally results in a shut-off of RNA transcription from DNA, while antisense RNA hybridization blocks translation of an mRNA molecule into polypeptide. Both techniques have been demonstrated to be effective in model systems. Information contained in the sequences of the present invention can be used to design antisense and triple helix-forming oligonucleotides, and other DNA binding agents.

6. Pharmaceutical Compositions and Vaccines

The present invention further provides pharmaceutical agents which can be used to modulate the growth or pathogenicity of *Borrelia burgdorferi*, or another related organism, *in vivo* or *in vitro*. As used herein, a "pharmaceutical agent" is defined as a composition of matter which can be formulated using known techniques to provide a pharmaceutical compositions. As used herein, the "pharmaceutical agents of the present invention" refers the pharmaceutical agents which are derived from the proteins encoded by the ORFs of the present invention or are agents which are identified using the herein described assays.

As used herein, a pharmaceutical agent is said to "modulate the growth pathogenicity of Borrelia burgdorferi or a related organism, in vivo or in vitro," when the agent reduces the rate of growth, rate of division, or viability of the organism in question. The pharmaceutical agents of the present invention can modulate the growth or pathogenicity of an organism in many fashions, although an understanding of the underlying mechanism of action is not needed to practice the use of the pharmaceutical agents of the present invention. Some agents will modulate the growth by binding to an important protein thus blocking the biological activity of the protein, while other agents may bind to a component of the outer surface of the organism blocking attachment or

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rendering the organism more prone to act the bodies nature immune system. Alternatively, the agent may comprise a protein encoded by one of the ORFs of the present invention and serve as a vaccine. The development and use of a vaccine based on outer membrane components are well known in the art.

As used herein, a "related organism" is a broad term which refers to any organism whose growth can be modulated by one of the pharmaceutical agents of the present invention. In general, such an organism will contain a homolog of the protein which is the target of the pharmaceutical agent or the protein used as a vaccine. As such, related organisms do not need to be bacterial but may be fungal or viral pathogens.

The pharmaceutical agents and compositions of the present invention may be administered in a convenient manner, such as by the oral, topical, intravenous, intraperitoneal, intramuscular, subcutaneous, intranasal or intradermal routes. The pharmaceutical compositions are administered in an amount which is effective for treating and/or prophylaxis of the specific indication. In general, they are administered in an amount of at least about 1 mg/kg body weight and in most cases they will be administered in an amount not in excess of about 1 g/kg body weight per day. In most cases, the dosage is from about 0.1 mg/kg to about 10 g/kg body weight daily, taking into account the routes of administration, symptoms, etc.

The agents of the present invention can be used in native form or can be modified to form a chemical derivative. As used herein, a molecule is said to be a "chemical derivative" of another molecule when it contains additional chemical moieties not normally a part of the molecule. Such moieties may improve the molecule's solubility, absorption, biological half life, *etc*. The moieties may alternatively decrease the toxicity of the molecule, eliminate or attenuate any undesirable side effect of the molecule, *etc*. Moieties capable of mediating such effects are disclosed in, among other sources, REMINGTON'S PHARMACEUTICAL SCIENCES (1980) cited elsewhere herein.

For example, such moieties may change an immunological character of the functional derivative, such as affinity for a given antibody. Such changes in immunomodulation activity are measured by the appropriate assay, such as a competitive type immunoassay. Modifications of such protein properties as redox or thermal stability, biological half-life, hydrophobicity, susceptibility to proteolytic degradation or the tendency to aggregate with carriers or into multimers also may be effected in this way and can be assayed by methods well known to the skilled artisan.

The therapeutic effects of the agents of the present invention may be obtained by providing the agent to a patient by any suitable means (e.g., inhalation, intravenously, intramuscularly, subcutaneously, enterally, or parenterally). It is preferred to administer the agent of the present invention so as to achieve an effective concentration within the blood or tissue in which the growth of the organism is to be controlled. To achieve an effective blood concentration, the preferred method is to administer the agent by injection. The administration may be by continuous infusion, or by single or multiple injections.

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In providing a patient with one of the agents of the present invention, the dosage of the administered agent will vary depending upon such factors as the patient's age, weight, height, sex, general medical condition, previous medical history, etc. In general, it is desirable to provide the recipient with a dosage of agent which is in the range of from about 1 pg/kg to 10 mg/kg (body weight of patient), although a lower or higher dosage may be administered. The therapeutically effective dose can be lowered by using combinations of the agents of the present invention or another agent.

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As used herein, two or more compounds or agents are said to be administered "in combination" with each other when either (1) the physiological effects of each compound, or (2) the serum concentrations of each compound can be measured at the same time. The composition of the present invention can be administered concurrently with, prior to, or following the administration of the other agent.

The agents of the present invention are intended to be provided to recipient subjects in an amount sufficient to decrease the rate of growth (as defined above) of the target organism.

The administration of the agent(s) of the invention may be for either a "prophylactic" or "therapeutic" purpose. When provided prophylactically, the agent(s) are provided in advance of any symptoms indicative of the organisms growth. The prophylactic administration of the agent(s) serves to prevent, attenuate, or decrease the rate of onset of any subsequent infection. When provided therapeutically, the agent(s) are provided at (or shortly after) the onset of an indication of infection. The therapeutic administration of the compound(s) serves to attenuate the pathological symptoms of the infection and to increase the rate of recovery.

The agents of the present invention are administered to a subject, such as a mammal, or a patient, in a pharmaceutically acceptable form and in a therapeutically effective concentration. A composition is said to be "pharmacologically acceptable" if its administration can be tolerated by a recipient patient. Such an agent is said to be administered in a "therapeutically effective amount" if the amount administered is physiologically significant. An agent is physiologically significant if its presence results in a detectable change in the physiology of a recipient patient.

The agents of the present invention can be formulated according to known methods to prepare pharmaceutically useful compositions, whereby these materials, or their functional derivatives, are combined in a mixture with a pharmaceutically acceptable carrier vehicle. Suitable vehicles and their formulation, inclusive of other human proteins, *e.g.*, human serum albumin, are described, for example, in REMINGTON'S PHARMACEUTICAL SCIENCES, 16th Ed., Osol, A., Ed., Mack Publishing, Easton PA (1980). In order to form a pharmaceutically acceptable composition suitable for effective administration, such compositions will contain an effective amount of one or more of the agents of the present invention, together with a suitable amount of carrier vehicle.

Additional pharmaceutical methods may be employed to control the duration of action. Control release preparations may be achieved through the use of polymers to complex or absorb one or more of the agents of the present invention. The controlled delivery may be effectuated by

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a variety of well known techniques, including formulation with macromolecules such as, for example, polyesters, polyamino acids, polyvinyl, pyrrolidone, ethylenevinylacetate, methylcellulose, carboxymethylcellulose, or protamine, sulfate, adjusting the concentration of the macromolecules and the agent in the formulation, and by appropriate use of methods of incorporation, which can be manipulated to effectuate a desired time course of release. Another possible method to control the duration of action by controlled release preparations is to incorporate agents of the present invention into particles of a polymeric material such as polyesters, polyamino acids, hydrogels, poly(lactic acid) or ethylene vinylacetate copolymers. Alternatively, instead of incorporating these agents into polymeric particles, it is possible to entrap these materials in microcapsules prepared, for example, by coacervation techniques or by interfacial polymerization with, for example, hydroxymethylcellulose or gelatine-microcapsules and poly(methylmethacylate) microcapsules, respectively, or in colloidal drug delivery systems, for example, liposomes, albumin microspheres, microemulsions, nanoparticles, and nanocapsules or in macroemulsions. Such techniques are disclosed in REMINGTON'S PHARMACEUTICAL SCIENCES (1980).

The invention further provides a pharmaceutical pack or kit comprising one or more containers filled with one or more of the ingredients of the pharmaceutical compositions of the invention. Associated with such container(s) can be a notice in the form prescribed by a governmental agency regulating the manufacture, use or sale of pharmaceuticals or biological products, which notice reflects approval by the agency of manufacture, use or sale for human administration.

In addition, the agents of the present invention may be employed in conjunction with other therapeutic compounds.

7. Shot-Gun Approach to Megabase DNA Sequencing

The present invention further demonstrates that a large sequence can be sequenced using a random shotgun approach. This procedure, described in detail in the examples that follow, has eliminated the up front cost of isolating and ordering overlapping or contiguous subclones prior to the start of the sequencing protocols.

Certain aspects of the present invention are described in greater detail in the examples that follow. The examples are provided by way of illustration. Other aspects and embodiments of the present invention are contemplated by the inventors, as will be clear to those of skill in the art from reading the present disclosure.

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ILLUSTRATIVE EXAMPLES

LIBRARIES AND SEQUENCING

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1. Shotgun Sequencing Probability Analysis

The overall strategy for a shotgun approach to whole genome sequencing follows from the Lander and Waterman (Landerman and Waterman, *Genomics* 2:231 (1988)) application of the equation for the Poisson distribution. According to this treatment, the probability, P0, that any given base in a sequence of size L, in nucleotides, is not sequenced after a certain amount, n, in nucleotides, of random sequence has been determined can be calculated by the equation P0 = e-m, where m is L/n, the fold coverage. For instance, for a genome of 2.8 Mb, m=1 when 2.8 Mb of sequence has been randomly generated (1X coverage). At that point, P0 = e-1 = 0.37. The probability that any given base has not been sequenced is the same as the probability that any region of the whole sequence L has not been determined and, therefore, is equivalent to the fraction of the whole sequence that has yet to be determined. Thus, at one-fold coverage, approximately 37% of a polynucleotide of size L, in nucleotides has not been sequenced. When 14 Mb of sequence has been generated, coverage is 5X for a 2.8 Mb and the unsequenced fraction drops to .0067 or 0.67%. 5X coverage of a 2.8 Mb sequence can be attained by sequencing approximately 17,000 random clones from both insert ends with an average sequence read length of 410 bp.

Similarly, the total gap length, G, is determined by the equation G = Le-m, and the average gap size, g, follows the equation, g = L/n. Thus, 5X coverage leaves about 240 gaps averaging about 82 bp in size in a sequence of a polynucleotide 2.8 Mb long.

The treatment above is essentially that of Lander and Waterman, *Genomics* 2: 231 (1988).

2. Random Library Construction

In order to approximate the random model described above during actual sequencing, a nearly ideal library of cloned genomic fragments is required. The following library construction procedure was developed to achieve this end.

Borrelia burgdorferi DNA is prepared by phenol extraction. A mixture containing 200 μ g DNA in 1.0 ml of 300 mM sodium acetate, 10 mM Tris-HCl, 1 mM Na-EDTA, 50% glycerol is processed through a nebulizer (IPI Medical Products) with a stream of nitrogen adjusted to 35 Kpa for 2 minutes. The sonicated DNA is ethanol precipitated and redissolved in 500 μ l TE buffer.

To create blunt-ends, a 100 μ l aliquot of the resuspended DNA is digested with 5 units of BAL31 nuclease (New England BioLabs) for 10 min at 30°C in 200 μ l BAL31 buffer. The digested DNA is phenol-extracted, ethanol-precipitated, redissolved in 100 μ l TE buffer, and then size-fractionated by electrophoresis through a 1.0% low melting temperature agarose gel. The section containing DNA fragments 1.6-2.0 kb in size is excised from the gel, and the LGT agarose is melted and the resulting solution is extracted with phenol to separate the agarose from the DNA. DNA is ethanol precipitated and redissolved in 20 μ l of TE buffer for ligation to vector.

A two-step ligation procedure is used to produce a plasmid library with 97% inserts, of which >99% were single inserts. The first ligation mixture (50 ul) contains 2 μg of DNA fragments, 2 µg pUC18 DNA (Pharmacia) cut with SmaI and dephosphorylated with bacterial alkaline phosphatase, and 10 units of T4 ligase (GIBCO/BRL) and is incubated at 14°C for 4 hr. The ligation mixture then is phenol extracted and ethanol precipitated, and the precipitated DNA is 5 dissolved in 20 µl TE buffer and electrophoresed on a 1.0% low melting agarose gel. Discrete bands in a ladder are visualized by ethidium bromide-staining and UV illumination and identified by size as insert (I), vector (v), v+I, v+2i, v+3i, etc. The portion of the gel containing v+I DNA is excised and the v+I DNA is recovered and resuspended into 20 μ l TE. The v+I DNA then is blunt-ended by T4 polymerase treatment for 5 min. at 37°C in a reaction mixture (50 ul) 10 containing the v+I linears, $500\,\mu\text{M}$ each of the 4 dNTPs, and 9 units of T4 polymerase (New England BioLabs), under recommended buffer conditions. After phenol extraction and ethanol precipitation the repaired v+I linears are dissolved in 20 µl TE. The final ligation to produce circles is carried out in a 50 μ l reaction containing 5 μ l of v+I linears and 5 units of T4 ligase at 14°C overnight. After 10 min. at 70°C the following day, the reaction mixture is stored at -20°C. 15

This two-stage procedure results in a molecularly random collection of single-insert plasmid recombinants with minimal contamination from double-insert chimeras (<1%) or free vector (<3%).

Since deviation from randomness can arise from propagation the DNA in the host, *E. coli* host cells deficient in all recombination and restriction functions (A. Greener, *Strategies 3 (1)*:5 (1990)) are used to prevent rearrangements, deletions, and loss of clones by restriction. Furthermore, transformed cells are plated directly on antibiotic diffusion plates to avoid the usual broth recovery phase which allows multiplication and selection of the most rapidly growing cells.

Plating is carried out as follows. A $100\,\mu l$ aliquot of Epicurian Coli SURE II 25 Supercompetent Cells (Stratagene 200152) is thawed on ice and transferred to a chilled Falcon 2059 tube on ice. A 1.7 μ l aliquot of 1.42 M beta-mercaptoethanol is added to the aliquot of cells to a final concentration of 25 mM. Cells are incubated on ice for 10 min. A 1 μ l aliquot of the final ligation is added to the cells and incubated on ice for 30 min. The cells are heat pulsed for 30 sec. at 42°C and placed back on ice for 2 min. The outgrowth period in liquid culture is eliminated from this protocol in order to minimize the preferential growth of any given 30 transformed cell. Instead the transformation mixture is plated directly on a nutrient rich SOB plate containing a 5 ml bottom layer of SOB agar (5% SOB agar: 20 g tryptone, 5 g yeast extract, 0.5 g NaCl, 1.5% Difco Agar per liter of media). The 5 ml bottom layer is supplemented with 0.4 ml of 50 mg/ml ampicillin per 100 ml SOB agar. The 15 ml top layer of SOB agar is supplemented with 1 ml X-Gal (2%), 1 ml MgCl2 (1 M), and 1 ml MgSO4/100 ml SOB agar. 35 The 15 ml top layer is poured just prior to plating. Our titer is approximately 100 colonies/10 µl aliquot of transformation.

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All colonies are picked for template preparation regardless of size. Thus, only clones lost due to "poison" DNA or deleterious gene products are deleted from the library, resulting in a slight increase in gap number over that expected.

3. Random DNA Sequencing

High quality double stranded DNA plasmid templates are prepared using a "boiling bead" method developed in collaboration with Advanced Genetic Technology Corp. (Gaithersburg, MD) (Adams *et al.*, *Science 252*:1651 (1991); Adams *et al.*, *Nature 355*:632 (1992)). Plasmid preparation is performed in a 96-well format for all stages of DNA preparation from bacterial growth through final DNA purification. Template concentration is determined using Hoechst Dye and a Millipore Cytofluor. DNA concentrations are not adjusted, but low-yielding templates are identified where possible and not sequenced.

Templates are also prepared from two Borrelia burgdorferi lambda genomic libraries. An amplified library is constructed in the vector Lambda GEM-12 (Promega) and an unamplified library is constructed in Lambda DASH II (Stratagene). In particular, for the unamplified lambda library, Borrelia burgdorferi DNA (> 100 kb) is partially digested in a reaction mixture (200 ul) containing 50 µg DNA, 1X Sau3AI buffer, 20 units Sau3AI for 6 min. at 23°C. The digested DNA was phenol-extracted and electrophoresed on a 0.5% low melting agarose gel at 2V/cm for 7 hours. Fragments from 15 to 25 kb are excised and recovered in a final volume of 6 ul. One μl of fragments is used with 1 μl of DASHII vector (Stratagene) in the recommended ligation reaction. One µl of the ligation mixture is used per packaging reaction following the recommended protocol with the Gigapack II XL Packaging Extract (Stratagene, #227711). Phage are plated directly without amplification from the packaging mixture (after dilution with $500\,\mu l$ of recommended SM buffer and chloroform treatment). Yield is about $2.5x103\,pfu/ul$. The amplified library is prepared essentially as above except the lambda GEM-12 vector is used. After packaging, about 3.5x104 pfu are plated on the restrictive NM539 host. The lysate is harvested in 2 ml of SM buffer and stored frozen in 7% dimethylsulfoxide. The phage titer is approximately 1x109 pfu/ml.

Liquid lysates (100 μ l) are prepared from randomly selected plaques (from the unamplified library) and template is prepared by long-range PCR using T7 and T3 vector-specific primers.

Sequencing reactions are carried out on plasmid and/or PCR templates using the AB Catalyst LabStation with Applied Biosystems PRISM Ready Reaction Dye Primer Cycle Sequencing Kits for the M13 forward (M13-21) and the M13 reverse (M13RP1) primers (Adams et al., Nature 368:474 (1994)). Dye terminator sequencing reactions are carried out on the lambda templates on a Perkin-Elmer 9600 Thermocycler using the Applied Biosystems Ready Reaction Dye Terminator Cycle Sequencing kits. T7 and SP6 primers are used to sequence the ends of the inserts from the Lambda GEM-12 library and T7 and T3 primers are used to sequence the ends of the inserts from the Lambda DASH II library. Sequencing reactions are performed

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by eight individuals using an average of fourteen AB 373 DNA Sequencers per day. All sequencing reactions are analyzed using the Stretch modification of the AB 373, primarily using a 34 cm well-to-read distance. The overall sequencing success rate very approximately is about 85% for M13-21 and M13RP1 sequences and 65% for dye-terminator reactions. The average usable read length is 485 bp for M13-21 sequences, 445bp for M13RP1 sequences, and 375 bp for dye-terminator reactions.

Richards *et al.*, Chapter 28 in AUTOMATED DNA SEQUENCING AND ANALYSIS, M. D. Adams, C. Fields, J. C. Venter, Eds., Academic Press, London, (1994) described the value of using sequence from both ends of sequencing templates to facilitate ordering of contigs in shotgun assembly projects of lambda and cosmid clones. We balance the desirability of bothend sequencing (including the reduced cost of lower total number of templates) against shorter read-lengths for sequencing reactions performed with the M13RP1 (reverse) primer compared to the M13-21 (forward) primer. Approximately one-half of the templates are sequenced from both ends. Random reverse sequencing reactions are done based on successful forward sequencing reactions. Some M13RP1 sequences are obtained in a semi-directed fashion: M13-21: sequences pointing outward at the ends of contigs are chosen for M13RP1 sequencing in an effort to specifically order contigs.

4. Protocol for Automated Cycle Sequencing

The sequencing is carried out using ABI Catalyst robots and AB 373 Automated DNA Sequencers. The Catalyst robot is a publicly available sophisticated pipetting and temperature control robot which has been developed specifically for DNA sequencing reactions. The Catalyst combines pre-aliquoted templates and reaction mixes consisting of deoxy- and dideoxynucleotides, the thermostable Taq DNA polymerase, fluorescently-labelled sequencing primers, and reaction buffer. Reaction mixes and templates are combined in the wells of an aluminum 96-well thermocycling plate. Thirty consecutive cycles of linear amplification (i.e.., one primer synthesis) steps are performed including denaturation, annealing of primer and template, and extension; i.e., DNA synthesis. A heated lid with rubber gaskets on the thermocycling plate prevents evaporation without the need for an oil overlay.

Two sequencing protocols are used: one for dye-labelled primers and a second for dye-labelled dideoxy chain terminators. The shotgun sequencing involves use of four dye-labelled sequencing primers, one for each of the four terminator nucleotide. Each dye-primer is labelled with a different fluorescent dye, permitting the four individual reactions to be combined into one lane of the 373 DNA Sequencer for electrophoresis, detection, and base-calling. ABI currently supplies pre-mixed reaction mixes in bulk packages containing all the necessary non-template reagents for sequencing. Sequencing can be done with both plasmid and PCR- generated templates with both dye-primers and dye- terminators with approximately equal fidelity, although plasmid templates generally give longer usable sequences.

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Thirty-two reactions are loaded per AB373 Sequencer each day, for a total of 960 samples. Electrophoresis is run overnight following the manufacturer's protocols, and the data is collected for twelve hours. Following electrophoresis and fluorescence detection, the ABI 373 performs automatic lane tracking and base-calling. The lane-tracking is confirmed visually. Each sequence electropherogram (or fluorescence lane trace) is inspected visually and assessed for quality. Trailing sequences of low quality are removed and the sequence itself is loaded via software to a Sybase database (archived daily to 8mm tape). Leading vector polylinker sequence is removed automatically by a software program. Average edited lengths of sequences from the standard ABI 373 are around 400 bp and depend mostly on the quality of the template used for the sequencing reaction. ABI 373 Sequencers converted to Stretch Liners provide a longer electrophoresis path prior to fluorescence detection and increase the average number of usable bases to 500-600 bp.

INFORMATICS

1. Data Management

A number of information management systems for a large-scale sequencing lab have been developed. (For review see, for instance, Kerlavage et al., Proceedings of the Twenty-Sixth Annual Hawaii International Conference on System Sciences, IEEE Computer Society Press, Washington D. C., 585 (1993)) The system used to collect and assemble the sequence data was developed using the Sybase relational database management system and was designed to automate data flow wherever possible and to reduce user error. The database stores and correlates all information collected during the entire operation from template preparation to final analysis of the genome. Because the raw output of the ABI 373 Sequencers was based on a Macintosh platform and the data management system chosen was based on a Unix platform, it was necessary to design and implement a variety of multi- user, client-server applications which allow the raw data as well as analysis results to flow seamlessly into the database with a minimum of user effort.

2. Assembly

An assembly engine (TIGR Assembler) developed for the rapid and accurate assembly of thousands of sequence fragments was employed to generate contigs. The TIGR assembler simultaneously clusters and assembles fragments of the genome. In order to obtain the speed necessary to assemble more than 104 fragments, the algorithm builds a hash table of 12 bp oligonucleotide subsequences to generate a list of potential sequence fragment overlaps. The number of potential overlaps for each fragment determines which fragments are likely to fall into repetitive elements. Beginning with a single seed sequence fragment, TIGR Assembler extends the current contig by attempting to add the best matching fragment based on oligonucleotide content. The contig and candidate fragment are aligned using a modified version of the Smith-Waterman algorithm which provides for optimal gapped alignments (Waterman, M. S., Methods

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in Enzymology 164:765 (1988)). The contig is extended by the fragment only if strict criteria for the quality of the match are met. The match criteria include the minimum length of overlap, the maximum length of an unmatched end, and the minimum percentage match. These criteria are automatically lowered by the algorithm in regions of minimal coverage and raised in regions with a possible repetitive element. The number of potential overlaps for each fragment determines which fragments are likely to fall into repetitive elements. Fragments representing the boundaries of repetitive elements and potentially chimeric fragments are often rejected based on partial mismatches at the ends of alignments and excluded from the current contig. TIGR Assembler is designed to take advantage of clone size information coupled with sequencing from both ends of each template. It enforces the constraint that sequence fragments from two ends of the same template point toward one another in the contig and are located within a certain range of base pairs (definable for each clone based on the known clone size range for a given library). The process resulted in 155 contigs as represented by SEQ ID NOs:1-155.

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3. Identifying Genes

The predicted coding regions of the *Borrelia burgdorferi* genome were initially defined with the program GeneMark, which finds ORFs using a probabilistic classification technique. The predicted coding region sequences were used in searches against a database of all nucleotide sequences from GenBank (July, 1997), using the BLASTN search method to identify overlaps of 50 or more nucleotides with at least a 95% identity (using default parameters). Those ORFs with nucleotide sequence matches are shown in Table 1. The ORFs without such matches were translated to protein sequences and compared to a non-redundant database of known proteins generated by combining the Swiss-prot, PIR and GenPept databases. ORFs that matched a database protein with BLASTP probability less than or equal to 0.01 are shown in Table 2. The table also lists assigned functions based on the closest match in the databases. ORFs that did not match protein or nucleotide sequences in the databases at these levels are shown in Table 3.

ILLUSTRATIVE APPLICATIONS

1. Production of an Antibody to a Borrelia burgdorferi Protein

Substantially pure protein or polypeptide is isolated from the transfected or transformed cells using any one of the methods known in the art. The protein can also be produced in a recombinant prokaryotic expression system, such as *E. coli*, or can be chemically synthesized. Concentration of protein in the final preparation is adjusted, for example, by concentration on an Amicon filter device, to the level of a few micrograms/ml. Monoclonal or polyclonal antibody to the protein can then be prepared as follows.

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2. Monoclonal Antibody Production by Hybridoma Fusion

Monoclonal antibody to epitopes of any of the peptides identified and isolated as described can be prepared from murine hybridomas according to the classical method of Kohler, G. and Milstein, C., *Nature 256*:495 (1975) or modifications of the methods thereof. Briefly, a mouse is repetitively inoculated with a few micrograms of the selected protein over a period of a few weeks. The mouse is then sacrificed, and the antibody producing cells of the spleen isolated. The spleen cells are fused by means of polyethylene glycol with mouse myeloma cells, and the excess unfused cells destroyed by growth of the system on selective media comprising aminopterin (HAT media). The successfully fused cells are diluted and aliquots of the dilution placed in wells of a microtiter plate where growth of the culture is continued. Antibody-producing clones are identified by detection of antibody in the supernatant fluid of the wells by immunoassay procedures, such as ELISA, as originally described by Engvall, E., *Meth. Enzymol. 70:*419 (1980), and modified methods thereof. Selected positive clones can be expanded and their monoclonal antibody product harvested for use. Detailed procedures for monoclonal antibody production are described in Davis, L. *et al.*, *Basic Methods in Molecular Biology*, Elsevier, New York. Section 21-2 (1989).

3. Polyclonal Antibody Production by Immunization

Polyclonal antiserum containing antibodies to heterogenous epitopes of a single protein can be prepared by immunizing suitable animals with the expressed protein described above, which can be unmodified or modified to enhance immunogenicity. Effective polyclonal antibody production is affected by many factors related both to the antigen and the host species. For example, small molecules tend to be less immunogenic than others and may require the use of carriers and adjuvant. Also, host animals vary in response to site of inoculations and dose, with both inadequate or excessive doses of antigen resulting in low titer antisera. Small doses (ng level) of antigen administered at multiple intradermal sites appears to be most reliable. An effective immunization protocol for rabbits can be found in Vaitukaitis, J. et al., J. Clin. Endocrinol. Metab. 33:988-991 (1971).

Booster injections can be given at regular intervals, and antiserum harvested when antibody titer thereof, as determined semi-quantitatively, for example, by double immunodiffusion in agar against known concentrations of the antigen, begins to fall. See, for example, Ouchterlony, O. et al., Chap. 19 in: Handbook of Experimental Immunology, Wier, D., ed, Blackwell (1973). Plateau concentration of antibody is usually in the range of 0. 1 to 0. 2 mg/ml of serum (about 12M). Affinity of the antisera for the antigen is determined by preparing competitive binding curves, as described, for example, by Fisher, D., Chap. 42 in: Manual of Clinical Immunology, second edition, Rose and Friedman, eds., Amer. Soc. For Microbiology, Washington, D. C. (1980)

Antibody preparations prepared according to either protocol are useful in quantitative immunoassays which determine concentrations of antigen-bearing substances in biological

samples; they are also used semi- quantitatively or qualitatively to identify the presence of antigen in a biological sample. In addition, antibodies are useful in various animal models of pneumococcal disease as a means of evaluating the protein used to make the antibody as a potential vaccine target or as a means of evaluating the antibody as a potential immunotherapeutic or immunoprophylactic reagent.

4. Preparation of PCR Primers and Amplification of DNA

Various fragments of the *Borrelia burgdorferi* genome, such as those of Tables 1-6 and SEQ ID NOS: 1-155 can be used, in accordance with the present invention, to prepare PCR primers for a variety of uses. The PCR primers are preferably at least 15 bases, and more preferably at least 18 bases in length. When selecting a primer sequence, it is preferred that the primer pairs have approximately the same G/C ratio, so that melting temperatures are approximately the same. The PCR primers and amplified DNA of this Example find use in the Examples that follow.

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5. Isolation of a Selected DNA Clone From B. burgdorferi

Three approaches are used to isolate a *B. burgdorferi* clone comprising a polynucleotide of the present invention from any *B. burgdorferi* genomic DNA library. The *B. burgdorferi* strain B31PU has been deposited as a convienent source for obtaining a *B. burgdorferi* strain although a wide varity of strains *B. burgdorferi* strains can be used which are known in the art.

B. burgdorferi genomic DNA is prepared using the following method. A 20ml overnight bacterial culture grown in a rich medium (e.g., Trypticase Soy Broth, Brain Heart Infusion broth or Super broth), pelleted, ished two times with TES (30mM Tris-pH 8.0, 25mM EDTA, 50mM NaCl), and resuspended in 5ml high salt TES (2.5M NaCl). Lysostaphin is added to final concentration of approx 50ug/ml and the mixture is rotated slowly 1 hour at 37C to make protoplast cells. The solution is then placed in incubator (or place in a shaking water bath) and warmed to 55C. Five hundred micro liter of 20% sarcosyl in TES (final concentration 2%) is then added to lyse the cells. Next, guanidine HCl is added to a final concentration of 7M (3.69g in 5.5 ml). The mixture is swirled slowly at 55C for 60-90 min (solution should clear). A CsCl gradient is then set up in SW41 ultra clear tubes using 2.0ml 5.7M CsCl and overlaying with 2.85M CsCl. The gradient is carefully overlayed with the DNA-containing GuHCl solution. The gradient is spun at 30,000 rpm, 20C for 24 hr and the lower DNA band is collected. The volume is increased to 5 ml with TE buffer. The DNA is then treated with protease K (10 ug/ml) overnight at 37 C, and precipitated with ethanol. The precipitated DNA is resuspended in a desired buffer.

In the first method, a plasmid is directly isolated by screening a plasmid *B. burgdorferi* genomic DNA library using a polynucleotide probe corresponding to a polynucleotide of the present invention. Particularly, a specific polynucleotide with 30-40 nucleotides is synthesized using an Applied Biosystems DNA synthesizer according to the sequence reported. The

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oligonucleotide is labeled, for instance, with ³²P-γ-ATP using T4 polynucleotide kinase and purified according to routine methods. (*See, e.g.*, Maniatis et al., Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Press, Cold Spring, NY (1982).) The library is transformed into a suitable host, as indicated above (such as XL-1 Blue (Stratagene)) using techniques known to those of skill in the art. *See, e.g.*, Sambrook et al. MOLECULAR CLONING: A LABORATORY MANUAL (Cold Spring Harbor, N.Y. 2nd ed. 1989); Ausubel et al., CURRENT PROTOCALS IN MOLECULAR BIOLOGY (John Wiley and Sons, N.Y. 1989). The transformants are plated on 1.5% agar plates (containing the appropriate selection agent, e.g., ampicillin) to a density of about 150 transformants (colonies) per plate. These plates are screened using Nylon membranes according to routine methods for bacterial colony screening. *See, e.g.*, Sambrook et al. MOLECULAR CLONING: A LABORATORY MANUAL (Cold Spring Harbor, N.Y. 2nd ed. 1989); Ausubel et al., CURRENT PROTOCALS IN MOLECULAR BIOLOGY (John Wiley and Sons, N.Y. 1989) or other techniques known to those of skill in the art.

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Alternatively, two primers of 15-25 nucleotides derived from the 5' and 3' ends of a polynucleotide of SEQ ID NOS:1-155 are synthesized and used to amplify the desired DNA by PCR using a *B. burgdorferi* genomic DNA prep as a template. PCR is carried out under routine conditions, for instance, in 25 µl of reaction mixture with 0.5 ug of the above DNA template. A convenient reaction mixture is 1.5-5 mM MgCl₂, 0.01% (w/v) gelatin, 20 µM each of dATP, dCTP, dGTP, dTTP, 25 pmol of each primer and 0.25 Unit of Taq polymerase. Thirty five cycles of PCR (denaturation at 94°C for 1 min; annealing at 55°C for 1 min; elongation at 72°C for 1 min) are performed with a Perkin-Elmer Cetus automated thermal cycler. The amplified product is analyzed by agarose gel electrophoresis and the DNA band with expected molecular weight is excised and purified. The PCR product is verified to be the selected sequence by subcloning and sequencing the DNA product.

Finally, overlapping oligos of the DNA sequences of SEQ ID NOS:1-155 can be chemically synthesized and used to generate a nucleotide sequence of desired length using PCR methods known in the art.

6(a). Expression and Purification Borrelia polypeptides in E. coli

The bacterial expression vector pQE60 is used for bacterial expression of some of the polypeptide fragements of the present invention. (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). pQE60 encodes ampicillin antibiotic resistance ("Ampr") and contains a bacterial origin of replication ("ori"), an IPTG inducible promoter, a ribosome binding site ("RBS"), six codons encoding histidine residues that allow affinity purification using nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin (QIAGEN, Inc., supra) and suitable single restriction enzyme cleavage sites. These elements are arranged such that an inserted DNA fragment encoding a polypeptide expresses that polypeptide with the six His residues (i.e., a "6

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X His tag") covalently linked to the carboxyl terminus of that polypeptide.

The DNA sequence encoding the desired portion of a *B. burgdorferi* protein of the present invention is amplified from *B. burgdorferi* genomic DNA using PCR oligonucleotide primers which anneal to the 5' and 3' sequences coding for the portions of the *B. burgdorferi* polynucleotide shown in SEQ ID NOS:1-155. Additional nucleotides containing restriction sites to facilitate cloning in the pQE60 vector are added to the 5' and 3' sequences, respectively.

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For cloning the mature protein, the 5' primer has a sequence containing an appropriate restriction site followed by nucleotides of the amino terminal coding sequence of the desired *B. burgdorferi* polynucleotide sequence in SEQ ID NOS:1-155. One of ordinary skill in the art would appreciate that the point in the protein coding sequence where the 5' and 3' primers begin may be varied to amplify a DNA segment encoding any desired portion of the complete protein shorter or longer than the mature form. The 3' primer has a sequence containing an appropriate restriction site followed by nucleotides complementary to the 3' end of the polypeptide coding sequence of SEQ ID NOS:1-155, excluding a stop codon, with the coding sequence aligned with the restriction site so as to maintain its reading frame with that of the six His codons in the pQE60 vector.

The amplified *B. burgdorferi* DNA fragment and the vector pQE60 are digested with restriction enzymes which recognize the sites in the primers and the digested DNAs are then ligated together. The *B. burgdorferi* DNA is inserted into the restricted pQE60 vector in a manner which places the *B. burgdorferi* protein coding region downstream from the IPTG-inducible promoter and in-frame with an initiating AUG and the six histidine codons.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described by Sambrook et al., *supra*.. *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kanr"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for expressing a *B. burgdorferi* polypeptide, is available commercially (QIAGEN, Inc., *supra*). Transformants are identified by their ability to grow on LB agar plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O/N") in liquid culture in LB media supplemented with both ampicillin (100 μ g/ml) and kanamycin (25 μ g/ml). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an optical density at 600 nm ("OD600") of between 0.4 and 0.6. Isopropyl- β -D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the lac repressor sensitive promoter, by inactivating the lacI repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

The cells are then stirred for 3-4 hours at 4°C in 6M guanidine-HCl, pH 8. The cell

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debris is removed by centrifugation, and the supernatant containing the *B. burgdorferi* polypeptide is loaded onto a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (QIAGEN, Inc., *supra*). Proteins with a 6 x His tag bind to the Ni-NTA resin with high affinity are purified in a simple one-step procedure (for details see: The QIAexpressionist, 1995, QIAGEN, Inc., *supra*). Briefly the supernatant is loaded onto the column in 6 M guanidine-HCl, pH 8, the column is first washed with 10 volumes of 6 M guanidine-HCl, pH 8, then washed with 10 volumes of 6 M guanidine-HCl pH 6, and finally the *B. burgdorferi* polypeptide is eluted with 6 M guanidine-HCl, pH 5.

The purified protein is then renatured by dialyzing it against phosphate-buffered saline (PBS) or 50 mM Na-acetate, pH 6 buffer plus 200 mM NaCl. Alternatively, the protein could be successfully refolded while immobilized on the Ni-NTA column. The recommended conditions are as follows: renature using a linear 6M-1M urea gradient in 500 mM NaCl, 20% glycerol, 20 mM Tris/HCl pH 7.4, containing protease inhibitors. The renaturation should be performed over a period of 1.5 hours or more. After renaturation the proteins can be eluted by the addition of 250 mM immidazole. Immidazole is removed by a final dialyzing step against PBS or 50 mM sodium acetate pH 6 buffer plus 200 mM NaCl. The purified protein is stored at 4°C or frozen at -80°C.

The polypeptide of the present invention are also prepared using a non-denaturing protein purification method. For these polypeptides, the cell pellet from each liter of culture is resuspended in 25 mls of Lysis Buffer A at 4°C (Lysis Buffer A = 50 mM Na-phosphate, 300 mM NaCl, 10 mM 2-mercaptoethanol, 10% Glycerol, pH 7.5 with 1 tablet of Complete EDTA-free protease inhibitor cocktail (Boehringer Mannheim #1873580) per 50 ml of buffer). Absorbance at 550 nm is approximately 10-20 O.D./ml. The suspension is then put through three freeze/thaw cycles from -70°C (using a ethanol-dry ice bath) up to room temperature. The cells are lysed via sonication in short 10 sec bursts over 3 minutes at approximately 80W while kept on ice. The sonicated sample is then centrifuged at 15,000 RPM for 30 minutes at 4°C. The supernatant is passed through a column containing 1.0 ml of CL-4B resin to pre-clear the sample of any proteins that may bind to agarose non-specifically, and the flow-through fraction is collected.

The pre-cleared flow-through is applied to a nickel-nitrilo-tri-acetic acid ("Ni-NTA") affinity resin column (Quiagen, Inc., *supra*). Proteins with a 6 X His tag bind to the Ni-NTA resin with high affinity and can be purified in a simple one-step procedure. Briefly, the supernatant is loaded onto the column in Lysis Buffer A at 4°C, the column is first washed with 10 volumes of Lysis Buffer A until the A280 of the eluate returns to the baseline. Then, the column is washed with 5 volumes of 40 mM Imidazole (92% Lysis Buffer A / 8% Buffer B) (Buffer B = 50 mM Na-Phosphate, 300 mM NaCl, 10% Glycerol, 10 mM 2-mercaptoethanol, 500 mM Imidazole, pH of the final buffer should be 7.5). The protein is eluted off of the column with a series of increasing Imidazole solutions made by adjusting the ratios of Lysis Buffer A to Buffer B. Three different concentrations are used: 3 volumes of 75 mM Imidazole, 3 volumes of

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150 mM Imidazole, 5 volumes of 500 mM Imidazole. The fractions containing the purified protein are analyzed using 8 %, 10 % or 14% SDS-PAGE depending on the protein size. The purified protein is then dialyzed 2X against phosphate-buffered saline (PBS) in order to place it into an easily workable buffer. The purified protein is stored at 4°C or frozen at -80°.

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The following alternative method may be used to purify B. burgdorferi expressed in E coli when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at $4-10^{\circ}$ C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells are harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells are then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 x g for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 x g centrifugation for 15 min., the pellet is discarded and the B. burgdorferi polypeptide-containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

Following high speed centrifugation (30,000 x g) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded *B. burgdorferi* polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 mm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the B. burgdorferi polypeptide are then pooled and mixed with 4

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volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A₂₈₀ monitoring of the effluent. Fractions containing the *B. burgdorferi* polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant *B. burgdorferi* polypeptide exhibits greater than 95% purity after the above refolding and purification steps. No major contaminant bands are observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. The purified protein is also tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

6(b). Alternative Expression and Purification Borrelia polypeptides in E. coli

The vector pQE10 is alternatively used to clone and express some of the polypeptides of the present invention for use in the soft tissue and systemic infection models discussed below. The difference being such that an inserted DNA fragment encoding a polypeptide expresses that polypeptide with the six His residues (i.e., a "6 X His tag") covalently linked to the amino terminus of that polypeptide. The bacterial expression vector pQE10 (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311) was used in this example. The components of the pQE10 plasmid are arranged such that the inserted DNA sequence encoding a polypeptide of the present invention expresses the polypeptide with the six His residues (i.e., a "6 X His tag")) covalently linked to the amino terminus.

The DNA sequences encoding the desired portions of a polypeptide of SEQ ID NOS:1-155 were amplified using PCR oligonucleotide primers from genomic *B. burgdorferi* DNA. The PCR primers anneal to the nucleotide sequences encoding the desired amino acid sequence of a polypeptide of the present invention. Additional nucleotides containing restriction sites to facilitate cloning in the pQE10 vector were added to the 5' and 3' primer sequences, respectively.

For cloning a polypeptide of the present invention, the 5' and 3' primers were selected to amplify their respective nucleotide coding sequences. One of ordinary skill in the art would appreciate that the point in the protein coding sequence where the 5' and 3' primers begins may be varied to amplify a DNA segment encoding any desired portion of a polypeptide of the present invention. The 5' primer was designed so the coding sequence of the 6 X His tag is aligned with the restriction site so as to maintain its reading frame with that of *B. burgdorferi* polypeptide. The 3' was designed to include an stop codon. The amplified DNA fragment was then cloned, and the protein expressed, as described above for the pQE60 plasmid.

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The DNA sequences of SEQ ID NOS:1-155 encoding amino acid sequences may also be cloned and expressed as fusion proteins by a protocol similar to that described directly above, wherein the pET-32b(+) vector (Novagen, 601 Science Drive, Madison, WI 53711) is preferentially used in place of pQE10.

The above methods are not limited to the polypeptide fragements actually produced. The above method, like the methods below, can be used to produce either full length polypeptides or desired fragements therof.

6(c). Alternative Expression and Purification of Borrelia polypeptides in E. coli

The bacterial expression vector pQE60 is used for bacterial expression in this example (QIAGEN, Inc., 9259 Eton Avenue, Chatsworth, CA, 91311). However, in this example, the polypeptide coding sequence is inserted such that translation of the six His codons is prevented and, therefore, the polypeptide is produced with no 6 X His tag.

The DNA sequence encoding the desired portion of the *B. burgdorferi* amino acid sequence is amplified from an *B. burgdorferi* genomic DNA prep the deposited DNA clones using PCR oligonucleotide primers which anneal to the 5' and 3' nucleotide sequences corresponding to the desired portion of the *B. burgdorferi* polypeptides. Additional nucleotides containing restriction sites to facilitate cloning in the pQE60 vector are added to the 5' and 3' primer sequences.

For cloning a *B. burgdorferi* polypeptides of the present invention, 5' and 3' primers are selected to amplify their respective nucleotide coding sequences. One of ordinary skill in the art would appreciate that the point in the protein coding sequence where the 5' and 3' primers begin may be varied to amplify a DNA segment encoding any desired portion of a polypeptide of the present invention. The 3' and 5' primers contain appropriate restriction sites followed by nucleotides complementary to the 5' and 3' ends of the coding sequence respectively. The 3' primer is additionally designed to include an in-frame stop codon.

The amplified *B. burgdorferi* DNA fragments and the vector pQE60 are digested with restriction enzymes recognizing the sites in the primers and the digested DNAs are then ligated together. Insertion of the *B. burgdorferi* DNA into the restricted pQE60 vector places the *B. burgdorferi* protein coding region including its associated stop codon downstream from the IPTG-inducible promoter and in-frame with an initiating AUG. The associated stop codon prevents translation of the six histidine codons downstream of the insertion point.

The ligation mixture is transformed into competent *E. coli* cells using standard procedures such as those described by Sambrook et al. *E. coli* strain M15/rep4, containing multiple copies of the plasmid pREP4, which expresses the lac repressor and confers kanamycin resistance ("Kanr"), is used in carrying out the illustrative example described herein. This strain, which is only one of many that are suitable for expressing *B. burgdorferi* polypeptide, is available commercially (QIAGEN, Inc., *supra*). Transformants are identified by their ability to grow on

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LB plates in the presence of ampicillin and kanamycin. Plasmid DNA is isolated from resistant colonies and the identity of the cloned DNA confirmed by restriction analysis, PCR and DNA sequencing.

Clones containing the desired constructs are grown overnight ("O/N") in liquid culture in LB media supplemented with both ampicillin ($100\,\mu g/ml$) and kanamycin ($25\,\mu g/ml$). The O/N culture is used to inoculate a large culture, at a dilution of approximately 1:25 to 1:250. The cells are grown to an optical density at 600 nm ("OD600") of between 0.4 and 0.6. isopropyl-b-D-thiogalactopyranoside ("IPTG") is then added to a final concentration of 1 mM to induce transcription from the *lac* repressor sensitive promoter, by inactivating the lacI repressor. Cells subsequently are incubated further for 3 to 4 hours. Cells then are harvested by centrifugation.

To purify the *B. burgdorferi* polypeptide, the cells are then stirred for 3-4 hours at 4°C in 6M guanidine-HCl, pH 8. The cell debris is removed by centrifugation, and the supernatant containing the *B. burgdorferi* polypeptide is dialyzed against 50 mM Na-acetate buffer pH 6, supplemented with 200 mM NaCl. Alternatively, the protein can be successfully refolded by dialyzing it against 500 mM NaCl, 20% glycerol, 25 mM Tris/HCl pH 7.4, containing protease inhibitors. After renaturation the protein can be purified by ion exchange, hydrophobic interaction and size exclusion chromatography. Alternatively, an affinity chromatography step such as an antibody column can be used to obtain pure *B. burgdorferi* polypeptide. The purified protein is stored at 4°C or frozen at -80°C.

The following alternative method may be used to purify *B. burgdorferi* polypeptides expressed in *E coli* when it is present in the form of inclusion bodies. Unless otherwise specified, all of the following steps are conducted at 4-10°C.

Upon completion of the production phase of the *E. coli* fermentation, the cell culture is cooled to 4-10°C and the cells are harvested by continuous centrifugation at 15,000 rpm (Heraeus Sepatech). On the basis of the expected yield of protein per unit weight of cell paste and the amount of purified protein required, an appropriate amount of cell paste, by weight, is suspended in a buffer solution containing 100 mM Tris, 50 mM EDTA, pH 7.4. The cells are dispersed to a homogeneous suspension using a high shear mixer.

The cells ware then lysed by passing the solution through a microfluidizer (Microfuidics, Corp. or APV Gaulin, Inc.) twice at 4000-6000 psi. The homogenate is then mixed with NaCl solution to a final concentration of 0.5 M NaCl, followed by centrifugation at 7000 x g for 15 min. The resultant pellet is washed again using 0.5M NaCl, 100 mM Tris, 50 mM EDTA, pH 7.4.

The resulting washed inclusion bodies are solubilized with 1.5 M guanidine hydrochloride (GuHCl) for 2-4 hours. After 7000 x g centrifugation for 15 min., the pellet is discarded and the B. burgdorferi polypeptide-containing supernatant is incubated at 4°C overnight to allow further GuHCl extraction.

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Following high speed centrifugation (30,000 x g) to remove insoluble particles, the GuHCl solubilized protein is refolded by quickly mixing the GuHCl extract with 20 volumes of buffer containing 50 mM sodium, pH 4.5, 150 mM NaCl, 2 mM EDTA by vigorous stirring. The refolded diluted protein solution is kept at 4°C without mixing for 12 hours prior to further purification steps.

To clarify the refolded *B. burgdorferi* polypeptide solution, a previously prepared tangential filtration unit equipped with 0.16 µm membrane filter with appropriate surface area (e.g., Filtron), equilibrated with 40 mM sodium acetate, pH 6.0 is employed. The filtered sample is loaded onto a cation exchange resin (e.g., Poros HS-50, Perseptive Biosystems). The column is washed with 40 mM sodium acetate, pH 6.0 and eluted with 250 mM, 500 mM, 1000 mM, and 1500 mM NaCl in the same buffer, in a stepwise manner. The absorbance at 280 mm of the effluent is continuously monitored. Fractions are collected and further analyzed by SDS-PAGE.

Fractions containing the *B. burgdorferi* polypeptide are then pooled and mixed with 4 volumes of water. The diluted sample is then loaded onto a previously prepared set of tandem columns of strong anion (Poros HQ-50, Perseptive Biosystems) and weak anion (Poros CM-20, Perseptive Biosystems) exchange resins. The columns are equilibrated with 40 mM sodium acetate, pH 6.0. Both columns are washed with 40 mM sodium acetate, pH 6.0, 200 mM NaCl. The CM-20 column is then eluted using a 10 column volume linear gradient ranging from 0.2 M NaCl, 50 mM sodium acetate, pH 6.0 to 1.0 M NaCl, 50 mM sodium acetate, pH 6.5. Fractions are collected under constant A₂₈₀ monitoring of the effluent. Fractions containing the *B. burgdorferi* polypeptide (determined, for instance, by 16% SDS-PAGE) are then pooled.

The resultant *B. burgdorferi* polypeptide exhibits greater than 95% purity after the above refolding and purification steps. No major contaminant bands are observed from Commassie blue stained 16% SDS-PAGE gel when 5 µg of purified protein is loaded. The purified protein is also tested for endotoxin/LPS contamination, and typically the LPS content is less than 0.1 ng/ml according to LAL assays.

6(d). Cloning and Expression of B. burgdorferi in Other Bacteria

B. burgdorferi polypeptides can also be produced in: B. burgdorferi using the methods of S. Skinner et al., (1988) Mol. Microbiol. 2:289-297 or J. I. Moreno (1996) Protein Expr. Purif. 8(3):332-340; Lactobacillus using the methods of C. Rush et al., 1997 Appl. Microbiol. Biotechnol. 47(5):537-542; or in Bacillus subtilis using the methods Chang et al., U.S. Patent No. 4,952,508.

7. Cloning and Expression in COS Cells

A B. burgdorferi expression plasmid is made by cloning a portion of the DNA encoding a

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B. burgdorferi polypeptide into the expression vector pDNAI/Amp or pDNAIII (which can be obtained from Invitrogen, Inc.). The expression vector pDNAI/amp contains: (1) an E. coli origin of replication effective for propagation in E. coli and other prokaryotic cells; (2) an ampicillin resistance gene for selection of plasmid-containing prokaryotic cells; (3) an SV40 origin of replication for propagation in eukaryotic cells; (4) a CMV promoter, a polylinker, an SV40 intron; (5) several codons encoding a hemagglutinin fragment (i.e., an "HA" tag to facilitate purification) followed by a termination codon and polyadenylation signal arranged so that a DNA can be conveniently placed under expression control of the CMV promoter and operably linked to the SV40 intron and the polyadenylation signal by means of restriction sites in the polylinker. The HA tag corresponds to an epitope derived from the influenza hemagglutinin protein described by Wilson et al. 1984 Cell 37:767. The fusion of the HA tag to the target protein allows easy detection and recovery of the recombinant protein with an antibody that recognizes the HA epitope. pDNAIII contains, in addition, the selectable neomycin marker.

A DNA fragment encoding a *B. burgdorferi* polypeptide is cloned into the polylinker region of the vector so that recombinant protein expression is directed by the CMV promoter. The plasmid construction strategy is as follows. The DNA from a *B. burgdorferi* genomic DNA prep is amplified using primers that contain convenient restriction sites, much as described above for construction of vectors for expression of *B. burgdorferi* in *E. coli*. The 5' primer contains a Kozak sequence, an AUG start codon, and nucleotides of the 5' coding region of the *B. burgdorferi* polypeptide. The 3' primer, contains nucleotides complementary to the 3' coding sequence of the *B. burgdorferi* DNA, a stop codon, and a convenient restriction site.

The PCR amplified DNA fragment and the vector, pDNAI/Amp, are digested with appropriate restriction enzymes and then ligated. The ligation mixture is transformed into an appropriate *E. coli* strain such as SURE™ (Stratagene Cloning Systems, La Jolla, CA 92037), and the transformed culture is plated on ampicillin media plates which then are incubated to allow growth of ampicillin resistant colonies. Plasmid DNA is isolated from resistant colonies and examined by restriction analysis or other means for the presence of the fragment encoding the *B. burgdorferi* polypeptide

For expression of a recombinant *B. burgdorferi* polypeptide, COS cells are transfected with an expression vector, as described above, using DEAE-dextran, as described, for instance, by Sambrook et al. (*supra*). Cells are incubated under conditions for expression of *B. burgdorferi* by the vector.

Expression of the *B. burgdorferi*-HA fusion protein is detected by radiolabeling and immunoprecipitation, using methods described in, for example Harlow et al., *supra*.. To this end, two days after transfection, the cells are labeled by incubation in media containing ³⁵S-cysteine for 8 hours. The cells and the media are collected, and the cells are washed and the lysed with detergent-containing RIPA buffer: 150 mM NaCl, 1% NP-40, 0.1% SDS, 1% NP-40, 0.5% DOC, 50 mM TRIS, pH 7.5, as described by Wilson et al. (*supra*). Proteins are



precipitated from the cell lysate and from the culture media using an HA-specific monoclonal antibody. The precipitated proteins then are analyzed by SDS-PAGE and autoradiography. An expression product of the expected size is seen in the cell lysate, which is not seen in negative controls.

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8. Cloning and Expression in CHO Cells

The vector pC4 is used for the expression of *B. burgdorferi* polypeptide in this example. Plasmid pC4 is a derivative of the plasmid pSV2-dhfr (ATCC Accession No. 37146). The plasmid contains the mouse DHFR gene under control of the SV40 early promoter. Chinese hamster ovary cells or other cells lacking dihydrofolate activity that are transfected with these plasmids can be selected by growing the cells in a selective medium (alpha minus MEM, Life Technologies) supplemented with the chemotherapeutic agent methotrexate. The amplification of the DHFR genes in cells resistant to methotrexate (MTX) has been well documented. *See, e.g.*, Alt et al., 1978, J. Biol. Chem. 253:1357-1370; Hamlin et al., 1990, Biochem. et Biophys. Acta, 1097:107-143; Page et al., 1991, Biotechnology 9:64-68. Cells grown in increasing concentrations of MTX develop resistance to the drug by overproducing the target enzyme, DHFR, as a result of amplification of the DHFR gene. If a second gene is linked to the DHFR gene, it is usually co-amplified and over-expressed. It is known in the art that this approach may be used to develop cell lines carrying more than 1,000 copies of the amplified gene(s). Subsequently, when the methotrexate is withdrawn, cell lines are obtained which contain the amplified gene integrated into one or more chromosome(s) of the host cell.

Plasmid pC4 contains the strong promoter of the long terminal repeat (LTR) of the Rouse Sarcoma Virus, for expressing a polypeptide of interest, Cullen, et al. (1985) Mol. Cell. Biol. 5:438-447; plus a fragment isolated from the enhancer of the immediate early gene of human cytomegalovirus (CMV), Boshart, et al., 1985, Cell 41:521-530. Downstream of the promoter are the following single restriction enzyme cleavage sites that allow the integration of the genes: Bam HI, Xba I, and Asp 718. Behind these cloning sites the plasmid contains the 3' intron and polyadenylation site of the rat preproinsulin gene. Other high efficiency promoters can also be used for the expression, e.g., the human \(\beta\)-actin promoter, the SV40 early or late promoters or the long terminal repeats from other retroviruses, e.g., HIV and HTLVI. Clontech's Tet-Off and Tet-On gene expression systems and similar systems can be used to express the B. burgdorferi polypeptide in a regulated way in mammalian cells (Gossen et al., 1992, Proc. Natl. Acad. Sci. USA 89:5547-5551. For the polyadenylation of the mRNA other signals, e.g., from the human growth hormone or globin genes can be used as well. Stable cell lines carrying a gene of interest integrated into the chromosomes can also be selected upon co-transfection with a selectable marker such as gpt, G418 or hygromycin. It is advantageous to use more than one selectable marker in the beginning, e.g., G418 plus methotrexate.

The plasmid pC4 is digested with the restriction enzymes and then dephosphorylated using calf intestinal phosphates by procedures known in the art. The vector is then isolated from

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a 1% agarose gel. The DNA sequence encoding the *B. burgdorferi* polypeptide is amplified using PCR oligonucleotide primers corresponding to the 5' and 3' sequences of the desired portion of the gene. A 5' primer containing a restriction site, a Kozak sequence, an AUG start codon, and nucleotides of the 5' coding region of the *B. burgdorferi* polypeptide is synthesized and used. A 3' primer, containing a restriction site, stop codon, and nucleotides complementary to the 3' coding sequence of the *B. burgdorferi* polypeptides is synthesized and used. The amplified fragment is digested with the restriction endonucleases and then purified again on a 1% agarose gel. The isolated fragment and the dephosphorylated vector are then ligated with T4 DNA ligase. *E. coli* HB101 or XL-1 Blue cells are then transformed and bacteria are identified that contain the fragment inserted into plasmid pC4 using, for instance, restriction enzyme analysis.

Chinese hamster ovary cells lacking an active DHFR gene are used for transfection. Five μg of the expression plasmid pC4 is cotransfected with 0.5 μg of the plasmid pSVneo using a lipid-mediated transfection agent such as Lipofectin™ or LipofectAMINE.™ (LifeTechnologies Gaithersburg, MD). The plasmid pSV2-neo contains a dominant selectable marker, the neo gene 15 from Tn5 encoding an enzyme that confers resistance to a group of antibiotics including G418. The cells are seeded in alpha minus MEM supplemented with 1 mg/ml G418. After 2 days, the cells are trypsinized and seeded in hybridoma cloning plates (Greiner, Germany) in alpha minus MEM supplemented with 10, 25, or 50 ng/ml of methotrexate plus 1 mg/ml G418. After about 10-14 days single clones are trypsinized and then seeded in 6-well petri dishes or 10 ml flasks 20 using different concentrations of methotrexate (50 nM, 100 nM, 200 nM, 400 nM, 800 nM). Clones growing at the highest concentrations of methotrexate are then transferred to new 6-well plates containing even higher concentrations of methotrexate (1 μ M, 2 μ M, 5 μ M, 10 mM, 20 mM). The same procedure is repeated until clones are obtained which grow at a concentration of 25 100-200 µM. Expression of the desired gene product is analyzed, for instance, by SDS-PAGE and Western blot or by reversed phase HPLC analysis.

The disclosure of all publications (including patents, patent applications, journal articles, laboratory manuals, books, or other documents) cited herein are hereby incorporated by reference in their entireties.

The present invention is not to be limited in scope by the specific embodiments described herein, which are intended as single illustrations of individual aspects of the invention. Functionally equivalent methods and components are within the scope of the invention, in addition to those shown and described herein and will become apparant to those skilled in the art from the foregoing description and accompanying drawings. Such modifications are intended to fall within the scope of the appended claims.

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

TABLE 1.

Contig C	ORF	Start (nt)	Stop	match	match gene name	% sim	%
	92	100363	100184	gil500722	similar to entire extracellular domain of glycine receptors	100	1 aen t
					[Caenorhabditis elegans]	3	}
1	537		513608 ₈	gil47453	ribosomal protein S12 [Streptococcus pneumoniae]	92	85
1	283			_	ATP-dependent protease ATPase subunit [Synechocystis sp.]	89	75
-	847	,	799131	gil467373	ribosomal protein S18 [Bacillus subtilis]	86	69
-	78				ribosomal protein L27 (rpL27) [Haemophilus influenzae]	85	70
	732	687538	686753 gill	şil1591672	phosphate transport system ATP-binding protein [Methanococcus jannaschii]	84	65
1	788	739513	739232 g	gil142459	initiation factor 1 [Bacillus subtilis]	84	89
	096	901448	901780 gnIII 69	gnllPIDle2437	PIDle2437 ORF YGL149w [Saccharomyces cerevisiae]	84	89
	760	717009	715843 E	23028	orf 361; ranslated orf similarity to SW: RF1_SALTY peptide chain release factor 1 of Salmonella typhimurium [Coxiella burnetii]	83	09
1	115		115312g	95315	NADH dehydrogenase subunit [Digitalis grandiflora]	82	58
_	184	178954	176918 bbs		EF-G=elongation factor G [Thermotoga maritima, Peptide, 682 aal [Thermotoga maritima]	82	63
1	447	425980	425453 g	gil143804	Ndk [Bacillus subtilis]	82	56
1	201	194702	194103 gil5	30438	arabinose transport protein [Mycoplasma capricolum]	81	53
1	477	446671	445589 gil8	82454	fructose 1,6-bisphosphate aldolase [Escherichia coli]	81	61
1	601	569453	568650 g	49227		81	56
-	887		837224 gil1	237019	Srb [Bacillus subtilis]	81	52
1	688		839497 _g	54276	peptide chain release factor 2 [Salmonella typhimurium]	81	65
1	968	∞	845440 g	377823	aminopeptidase [Bacillus subtilis]	81	09
1	8		68890 gil 1	1619909	DNA mismatch repair protein [Thermotoga maritima]	80	59
1	354		349157 gill	165976	chemotaxis protein Che Y [Treponema pallidum]	08	42
	423	409238	408855 g	gnllPIDle2118 29	PIDIe2118 50S ribosomal protein L14 [Odontella sinensis]	08	61
I	426	410130	409711 _g		50S ribosomal protein L16 [Synechocystis sp.]	08	59
	507	482736	482936 g	15924	glucosyltransferase [Saccharomyces cerevisiae]	80	40
	534	505081	505467 _p	pirlA02771IR	ribosomal protein L7/L12 - Micrococcus luteus	80	67

		PCT	US98/	12764
54	56	09	38	47

		0 59	9 62	99 6	09		50		58		09	51	7 59			52	54	99	09	38	47		
		8	79	79	79	78	78	78	78	78	77	77	77	77	17	77	77	77	9/	76	9/	76	76
Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins	7MCML	gilS	9994 gnllPIDle2426 arginine deiminase [Clostridium perfringens]	446835 gnllPIDle2881 glucose epimerase [Bacillus thuringiensis]	757704 gil455176 glucosamine-6-phosphate deaminase protein [Escherichia coli]	17809	134323 gil159199 cecropin D [Hyalophora cecropia]	216028 gnllPIDle2655 DnaJ-homologue [Thermus aquaticus thermophilus]	503849 gil587583 ribosomal protein L11 [Thermus aquaticus thermophilus]		127745 gil537364 heat shock protein 60 (GroEL) like protein [Porphyromonas gingivalis]	5682	Phosphoglycerate mutase 1 [Escherichia co	1349	3746	531370 gil143795 [transfer RNA-Tyr synthetase [Bacillus subtilis]	892 gil 1653602 [hypothetical protein [Synechocystis sp.]	790909 gnllPIDle2488 unknown [Mycobacterium tuberculosis]	6	141736 bbs/77721 KHS toxin, killer heat sensitive toxin=KHS [Saccharomyces cerevisiae]		5	355508 aik33147 rihosa phosphata arranhosahaltingg [Docilling 11-1-1-1
	Ш			447926		31595	134667	215177 2	L.		~	182991		272770 2				790115 7	62205		Ш		356740 3
		597	9	478	804	25	134	230	531	298	127	190	225	284	324	555	770	833	52	144	293	323	363
				1	1		1		1	1	=		=		1	1	1	-	_	-		=	=

410	403332	Borrelia bur 402922 91	burgdorferi - Puta 91606232	rgdorferi - Putative coding regions of novel proteins similar to know proteins 606232 130S ribosomal subunit protein S11 (Escherichia coli)	76	52
403754	754	403341		30S ribosomal protein \$13 [Synechocystis sp.]	76	55
431743	743	431003	gil1016012	neural cell adhesion protein BIG-2 precursor [Rattus norvegicus]	9/	19
670457	457	671569 gil	gi 467376	unknown [Bacillus subtilis]	76	58
824	824849	826675	gil1303804	YqeQ [Bacillus subtilis]	16	52
886017	017	886751	gil1183839	unknown [Pseudomonas aeruginosa]	9/	54
5	9956	8943	gil1552842	OTCase [Escherichia coli]	75	62
9	61909	59735	gil1184680	polynucleotide phosphorylase [Bacillus subtilis]	75	54
9	66283	63620	gil39954	IF2 (aa 1-741) [Bacillus stearothermophilus]	75	53
6	93454	94410	gnllPIDle2891 38	similar to flagellar hook-basal body proteins [Bacillus subtilis]	75	46
	97435	98283	gil687583	RpoS [Yersinia enterocolitica]	75	47
2.	229112	230158	gil1574806	spermidine/putrescine transport ATP-binding protein (potA) [Haemophilus influenzae]	75	55
2	251076	250801	gil1763634	alpha1 A-voltage-dependent calcium channel [Homo sapiens]	75	09
2	285723	284461	gil556886	serine hydroxymethyltransferase [Bacillus subtilis]	75	58
6	367682	366903	gil467372	3'-exo-deoxyribonuclease [Bacillus subtilis]	75	62
(c.)	378055	377114	gil45986	NAD synthetase [Rhodobacter capsulatus]	75	55
4	406437	405925	gil1044981	ribosomal protein S5 [Bacillus subtilis]	75	99
4	407390		gi	L6 ribosomal protein [Streptomyces coelicolor]	75	53
4	409520	409251	gil44218	ribosomal protein S17 (AA 1-85) [Mycoplasma capricolum]	75	58
	502806	503366	gil396321	nusG [Escherichia coli]	75	99
'	523428	522904	gil1573470	H. influenzae predicted coding region H10491 [Haemophilus influenzae]	75	55
\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \	546579	548393	pirlC30010lC	hypothetical ORF-6 protein - Sauroleishmania tarentolae	75	50
			30010	mitochondrion (SGČ6)		
∞	854433	855215	gil511148	hemolysin [Serpulina hyodysenteriae]	75	99
	85054	83102	gil467458	cell division protein [Bacillus subtilis]	74	57
1	158608	157502gil5	gil531460	Mbl protein [Bacillus subtilis]	74	49
<u> </u>	172327	171950 pirl/ 454	pirlA45434lA 45434	ribosomal protein L19 - Bacillus stearothermophilus	74	54
44	443773	445203	gil396501	aspartyl-tRNA synthetase [Thermus aquaticus thermophilus]	74	52
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wo	98	3/58	943	3)					6	8							PO	C T /	US	98/	127	64
	57	58	49	50	56	49	43	53	51	51	52	53	55	53		47	54	56	63	20	53	54	51	49	51	51	41	42
	74	74	74	74	73	73	73	73	73	73	73	73	73	73		73	73	73	73	73	72	72	72	72	72	72	72	72
Putative coding regions of novel proteins similar to know proteins	S-adenosylmethionine synthetase [Staphylococcus aureus]	enolase [Bacillus subtilis]	hypothetical protein [Synechocystis sp.]	UDP-glucose pyrophosphorylase [Bacillus subtilis]	excinuclease ABC subunit A [Synechocystis sp.]	sensor kinase [Bacillus subtilis]	Erg8p [Saccharomyces cerevisiae]	'ORF' [Escherichia coli]	ORF YLR069c [Saccharomyces cerevisiae]	hypothetical [Haemophilus influenzae]	hemolysin [Serpulina hyodysenteriae]	glycoprotein 120 [Simian immunodeficiency virus]	sporulation protein [Bacillus subtilis]	60 kda antigen [Borrelia coriaceae, C053, ATCC 4338, Peptide,	[514 aa] [Borrelia coriaceae]	type-I signal peptidase SpsB [Staphylococcus aureus]	unknown [Mycobacterium tuberculosis]	Similar to Seryl-tRNA synthetase [Saccharomyces cerevisiae]	36 ORF YGR248w [Saccharomyces cerevisiae]	hypothetical protein [Synechocystis sp.]	hemolysin [Serpulina hyodysenteriae]	NtrC/NifA-like protein regulator [Escherichia coli]	Similar to Saccharomyces cerevisiae SUA5 protein [Bacillus subtilis]	transcription-repair coupling factor [Bacillus subtilis]	ribosomal protein S4 (rpS4) [Haemophilus influenzae]	lon protease [Bacillus brevis]	haemolysin releasing protein (AA 1-548) [Vibrio cholerae]	CTP synthase [Methanococcus jannaschii]
gdorferi -	gil1020317	564347 gil460259	680489 gil1651962	701173 gil289287	17551 gil1652531	104947 gil 514330	181102 gil887601	302786 gil473817	361078 gnilPIDle2457	423181 gil1574704	533672 gil511145	548045 gil406135	567504 gil 143607	570729 bbs1161785		gil1595810	651727 gnllPIDle2684 56	680499 gil500705	 2	844964 gil1652288	26497 gil511145	106305 gil619917	135055 gil556881	260308 gil467444	268221 gil1573812	268472 gil402504	318363 gil48362	321053 gil1591801
	524561	565672	681529	702297	20409	103790	182064	303616	358916	424047	531372	548257	568379	572375		634175	654267	679186	682189	845455	24242	104935	134036	256925	267529	270922	319544	322678
	549	595	720	745	13	86	188	314	366	444	556	276	865	604		674	692	719	725	895	91	66	133	270	280	282	325	328

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		23	55	54	51	47	48	54	52	42	43	48	52	48	54	52	52	42	53	56	53	32	50	58	41		48
C	/2/	72	72	72	72	72	72	72	71	71	71	71	71	711	71	71	71	71	71	71	70	70	70	70	70	-	70
	pism protein [Eschenchia con]	M. genitalium predicted coding region MG246 [Mycoplasma genitalium]	S14 protein (AA 1-61) [Bacillus subtilis]	ribosomal protein L13 (rpL13) [Haemophilus influenzae]	sporulation protein (spoIIIE) [Haemophilus influenzae]		large ribosomal subunit protein L35 [Buchnera aphidicola]	asparaginyl-tRNA synthetase [Synechocystis sp.]	UvrB [Helicobacter pylori]	beta-b protein [Barley stripe mosaic virus]	ORF9 [Rhizobium meliloti]			similar to multifunctional aminoacyl-tRNA synthetase, especially to the prolyl-tRNA synthetase region [Caenorhabditis elegans]		pyruvate kinase [Bacillus stearothermophilus]	ORF1 [Synechococcus elongatus]	secretion protein SecY (AA 1-482) [Mycoplasma capricolum]	ORF for methionine amino peptidase [Bacillus subtilis]	queA [Escherichia coli]	Cdc28p [Schizosaccharomyces pombe]	o287 [Escherichia coli]	[flgG protein product (AA 1-260) [Salmonella typhimurium]	H. influenzae predicted coding region HI1534 [Haemophilus influenzae]	(AE000012) Mycoplasma pneumoniae, phosphocarrier protein HPr; similar to GenBank Accession Number A49683, from M.	capricolum [Mycoplasma pneumoniae]	RL 50S RIBOSOMAL PROTEIN L1 (BL1).
burgdorferi -	gII14500/	gil 1045937	gil580930				50	60	50	gil1016781	gil534842	gil1652099	gil1732243	gil459009	530156 pirlS585221S5 8522	gil285623	gil217121	gil44228	_	gil	gil	gil290494	gil47677	gil1574387	gil1673757		splQ06797IRL
Borrelia	241102	399096	407779	563850	643399	710750	721640	810511	20407	87674	278239	296736	312130	496383	530156	552271	643661	681561	807700	890665	40613	44806	95220 gil	128569	441330		504529 sp
241460	341400	399941	408009	563383	641030	710160	721422	811923	22434	87471	278760	298685	313551	494911	528795	553725	644626	681731	806939	890096	38112	45750	94408	127889	441049		503834
340	040	405	420	593	682	754	191	098	14	72	289	307	321	225	554	582	684	723	856	947	28	36	84	128	468		532
	1					1	1		1		1	1	Ţ	I		1	I	1	-			_	T		1		

			Borrelia burgd	ourgdorferi - Puta	dorferi - Putative coding regions of novel proteins similar to know proteins		
	594	563858	564280 g	6169	30S ribosomal subunit protein S9 [Escherichia coli]	70	26
1	622	591070	591606 gill.	1153906	CheW protein [Salmonella typhimurium]	70	48
	703	664161	662611 g	PIDIe2839	glycerol kinase [Sulfolobus solfataricus]	70	09
	726	682886	682659 g	gil836815	cdc4 gene product which is essential for initiation of DNA replication in yeast [Saccharomyces cerevisiae]	0/	35
1	99/	720854	721417g	gil436165	Dsg [Myxococcus xanthus]	70	47
	892	721649	722008 g 8	gniiPIDle2549 81	PDle2549 ribosomal protein L20 [Bacillus subtilis]	70	48
	596	904395	905465 _g	100074	tryptophanyl-tRNA synthetase [Clostridium longisporum]	70	47
1	87	98696	97336g	60092	asparagine-rich protein [Plasmodium falciparum]	69	46
1	011	112658	113602 g	001733	ABC transporter [Synechocystis sp.]	69	46
	181	174037	173762 pirl(471	247154IC 54	ribosomal protein S16 - Bacillus subtilis	69	52
	233	219872	218076 gill	001493	protein-export membrane protein SecD [Synechocystis sp.]	69	47
1	234	220245	\sim		ORF11 [Enterococcus faecalis]	69	32
I	373	366148	_		hypothetical [Haemophilus influenzae]	69	48
	419	407781	Ī	98771	ribosomal S8 protein [Thermus aquaticus thermophilus]	69	46
1	517	489315	491207 gill	51932	fructose enzyme II [Rhodobacter capsulatus]	69	42
1	009	568891	568388 gil 14	3606	sporulation protein [Bacillus subtilis]	69	44
1	733	860689	687536 gil	03826	YqgI [Bacillus subtilis]	69	46
	874	826778	827746 pirlS 8183	08183180	L-lactate dehydrogenase (EC 1.1.1.27) X - Bacillus psychrosaccharolyticus	69	20
	894	844392		92324	M. jannaschii predicted coding region MJ1172 [Methanococcus jannaschii]	69	53
Ţ	934	879725	879237 g	53566	ORF (19K protein) [Enterococcus faecalis]	69	42
-	46	<i>61118</i>	57976 gil80	11809583	unknown [Saccharomyces cerevisiae]	89	36
-	107	110374	111513 gnllP 43	nIIPIDIe2559 3	PIDIe2559 M04B2.4 [Caenorhabditis elegans]	89	48
1	132	133978	133148 gil1	il1001663	rare lipoprotein A [Synechocystis sp.]	89	53
1	142	141239	142642 gnllPl 74	nIIPIDIe2338 4	'IDle2338 hypothetical protein [Bacillus subtilis]	89	45
1	148	145381	144005 gil55	8574	pyrophosphatefructose-6-phosphate 1-phosphotransferase	89	48

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	42	50	45	42	41	47	42	20	40	4	51	37	42	51	52	52	_		44	20	50	47	52	43	49	47
	89	89	89	89	89	89	89	89	89	89	89	89	89	89	89	89			89	89	89	89	89	29	29	19
Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins [HEntamoeba histolytica]	beta-galactoside binding protein [Mus musculus]	ORF3 [Bacillus subtilis]	signal recognition particle protein [Synechocystis sp.]	galactose binding protein [Escherichia coli]	hypothetical [Haemophilus influenzae]	glucose 6-phosphate dehydrogenase [Synechocystis sp.]	fructose-permease IIBC component (fruA) [Haemophilus influenzae]	MHc, class IIB gene product [Poecilia reticulata]	ORF 1 [Mycoplasma mycoides]	SecY protein [Corynebacterium glutamicum]	ribosomal protein L29 [Methanococcus jannaschii]	adenylate kinase [Paracoccus denitrificans]	'ORF' [Escherichia coli]	hypothetical protein [Bacillus subtilis]	D9461.18p; CAI: 0.15 [Saccharomyces cerevisiae]	coded for by C. elegans cDNA CEESS55F; coded for by C. elegans cDNA vk84a1 3; coded for by C. elegans cDNA	yk78g7.3; coded for by C. elegans cDNA yk168g9.5; coded for	by C. elegans cDNA yk/8g/.5; coded for by C. elegans cDNA yk84a1.5; strong s	outer surface protein F [Borrelia burgdorferi]	hypothetical protein [Synechocystis sp.]	glcA gene product [Staphylococcus carnosus]	ORF III [Escherichia coli]	IllPIDle2902 polypeptide deformylase [Calothrix PCC7601]	IIPIDIe2804 unknown [Streptococcus pneumoniae]	ILS1 protein [Saccharomyces cerevisiae]	DNA topoisomerase I [Synechocystis sp.]
orrelia burgdorferi - Put 	147295 gil193442	169296 gil1389549	50	ЬÒ	227344 gi 1573101	B	251077 gil1573422	363333 gil976104	399874 gil150209	امةا	409508 gil1591164	ಹ	524549 gil473817	657673 gnIIPIDIe2551 17	1460 gil927711	686750 gil1707057			gi	gil	797807 gil 1072418	827776 gil147774		24255 gnllPIDle2804 90	29640 gil498991	139 gil1652946
Bo	147107 147	170051 169					249185 251	364016 363	401421 399	L.		482737 482		658281 657	1	989 506589				_		2	850141 850	22531 24		35545 38
	151	173	182	203	243	255	263	372	407	412	425	90\$	055	L69	717	05/			747	793	843	875	006	15	20	27
				1	1	1							1							1						

777	7/6611	114333 gii1001329	001529 [hypothetical protein [Synechocystis sp.]	29	36
1 170	166286	gil5	CapE [Staphylococcus aureus]	19	35
1 202	195499	194651 gil1674275	(AE000056) Mycoplasma pneumoniae, hypothetical ABC transporter (yjcW) homolog; similar to Swiss-Prot Accession Number P32721, from E. coli [Mycoplasma pneumoniae]	<i>L</i> 9	41
1 206	197487	Bill	P protein [Synechocystis sp.]	<i>L</i> 9	35
1 271	260292	gil34	acetate kinase [Methanosarcina thermophila]	<i>L</i> 9	44
1 313	302731	301643 gnllPIDle2499 81	PIDle 2499 phosphotransacety lase [Thermoanaerobacterium thermosaccharolyticum]	<i>L</i> 9	51
1 422	408897	408535 pirlA02819IR 5BS24	ribosomal protein L24 - Bacillus stearothermophilus	19	49
1 480	450326	gill	hypothetical [Haemophilus influenzae]	<i>L</i> 9	42
1 529	502315	502509 gil1001264	50S ribosomal protein L33 [Synechocystis sp.]	19	56
1 588	559618	gill	amidase [Moraxella catarrhalis]	1.9	51
1 683	643676		ribosomal protein S21 [Myxococcus xanthus]	19	49
1 698	658454		TagE [Vibrio cholerae]	<i>L</i> 9	38
1 700	660039	660536 gil 467420	unknown [Bacillus subtilis]	19	42
1 729	684089	685888 gnllPIDIe2676 07	685888 gnllPIDle2676 alanyl-tRNA synthatase [Thermus aquaticus thermophilus]	<i>L</i> 9	51
1 835	791754	792341 gnllPIDle2487 63	gnllPIDle2487 unknown [Mycobacterium tuberculosis]	<i>L</i> 9	46
1 857	807722	gill:	GsrA protein [Yersinia enterocolitica]	19	46
1 868	819577	820905 gil1590954	ATP synthase, subunit B [Methanococcus jannaschii]	<i>L</i> 9	53
1 74	88393	88028 gil1572979	hypothetical [Haemophilus influenzae]	99	43
1 91	99152	100252 gil561690	sialoglycoprotease [Pasteurella haemolytica]	99	44
1 123	121472		endonuclease III [Synechocystis sp.]	99	42
1 149	146362	gil12	orf304 gene product [Treponema pallidum]	99	43
1 185	179585	gil15	neutrophil activating protein (napA) [Haemophilus influenzae]	99	49
1 275	265075	gil40	cytidine deaminase [Mycoplasma pirum]	99	41
1 330	324514	323696 gil1574641	ribonucleotide transport ATP-binding protein (mkl) [Haemophilus influenzae]	99	41
335	327265		che Y gene product [Rhodobacter sphaeroides]	99	44
1 355	349142	382	Flis (Bacillus enhtilis)		00

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53 53 41 40 40	33 47 47	33 46 40 40 39 42	41 43 47 47	4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4
99 99	99	99 65 65 65 65 65	65 65 65 65 65 65 65 65 65 65 65 65 65 6	65 65 65 65 65 65
Putative coding regions of novel proteins similar to know proteins [tar-1 [Trichostrongylus colubriformis] hemolysin [Serpulina hyodysenteriae] P. putida genes rpmH, rnpA, 9k, 60k, 50k, gidA, gidB, uncl and uncB [Pseudomonas putida] methyltransferase (cheR; EC 2.1.1.24) [Salmonella typhimurium] A 'c' was inserted after nt 369 (=nt 10459 in genomic sequence (M10126)) to correct -1 frameshift probably due to gel compression [Leishmania tarentolae]	putative pectinesterase [Medicago sativa] 9 OrfD [Streptococcus pneumoniae] could accelerate degradation of certain transcripts [Bacillus subtilis]	glycerol ester hydrolase [Staphylococcus aureus] novel hemolytic factor [Bacillus cereus] similar to the ATP-binding transport protein family [Buchnera aphidicola] spo VG gene product [Bacillus megaterium] phosphatidylserine decarboxylase [Bacillus subtilis] ClpP [Yersinia enterocolitica]	pencillin-oliding protein 2 (pbp2) [Haemophilus influenzae] poly(A) polymerase [Bacillus subtilis] bacterial cell wall hydrolase [Enterococcus faecalis] DNA ligase (lig) [Haemophilus influenzae] Pz-peptidase [Bacillus licheniformis] DNA mismatch repair protein [Aquifex pyrophilus]	single-stranded DNA-binding protein [Synechocystis sp.] gyrase A [Helicobacter pylori] leader peptidase I [Synechocystis sp.] YbbQ [Bacillus subtilis] hypothetical [Haemophilus influenzae] UDP-N-acetyl muramate-alanine ligase [Bacillus subtilis]
Borrelia burgdorferi - 350827 gil1546788 398324 gil296626 460550 gil45713 485159 gil153903 527316 gil340613	581069 gil886130 596288 gnllPIDle26 31 723522 gil1762342	770060 gil393266 795208 gil662880 834262 gil862629 87619 gil39656 102803 gil532272 109649 gil1377852	169323 169323 346553 655781	7988.27 gil1001362 878559 gil508471 882224 gil1652260 901519 gil1256146 904407 gil1573307 45683 gil556014
351051 399121 461335 486046 526495	595395 595395 723788	770251 795927 835002 87915 103039 110281	168084 255918 348568 657577 695297	876643 881238 902331 903280 47101
358 404 491 513 552	627 772	816 841 882 73 97 106	172 268 353 353 696 741	932 936 961 963 37

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44 43 45 45	347	33 33 4	44 41 45 45	42 42 41	38 38 38 41 41 45
2 4 4 4 4	2 2 2 2	2 2 2 2	46 64 46	64 64	63 64 66 66
- Putative coding regions of novel proteins similar to know proteins I rhoptry protein [Plasmodium yoelii] valyl-tRNA synthetase (valS) [Haemophilus influenzae] threonyl-tRNA synthetase (thrS; EC 6.1.1.3) [Escherichia coli] acyl carrier protein [Synechocystis sp.] lipopolysaccharide core biosynthesis protein (kdtB) [Haemophilus influenzae]	2488 unknown [Mycobacterium tuberculosis] ORF2136 [Marchantia polymorpha]	hypothetical protein (GB:U00021_5) [Mycoplasma genitalium] transmembrane protein [Escherichia coli] unknown [Bacillus subtilis] cheW peptide [Escherichia coli]	monophosphatase [Synechocystis sp.] DNA polymerase III subunit [Bacillus subtilis] protein-glutamate methylesterase (EC 3.1.1.61) - Salmonella typhimurium dipeptide transport system permease protein (dppB) [Haemophilus influenzae]	soluble lytic transglycosylase [Synechocystis sp.] unknown [Mycobacterium tuberculosis] W04B2.3 [Caenorhabditis elegans]	hypothetical [Haemophilus influenzae] glutamate synthase [Escherichia coli] v-type Na-ATPase [Enterococcus hirae] methionyl-tRNA formyltransferase [Escherichia coli] thioredoxin [Arabidopsis thaliana] SbcC (AA 1-1048) [Escherichia coli]
Borrelia burgdorferi 71642 gil104178 129336 gil157422 151140 gil43066 170033 gil165239 170545 gil157365	173513 gnllPIDle7 173513 gnllPIDle7 93 197436 gil11665 205761 gil165286			640224 710194 771969	795211 gil1573939 812853 gil396314 823339 gil472918 851615 gil581088 853884 gil992960 31444 gil42914
72211 131969 152924 170326 171105	173764 173764 197654 206795	228146 228146 230149 253160 333349	376509 428137 484558 570416	637996 709637 771784	793892 811972 821501 850668 853492 34314
130 130 174 175	207	244 246 267 340	384 449 510 603	679 753 817	839 . 861 870 901 904 24
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45	42	41	36	36	46	41	42	34	48	49	43	47	37	45	36	33	28	40	43	36	38	41	36	48	34
63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	63	62	62	62	62	62
urgdorferi - Putative coding regions of novel proteins similar to know proteins II1652022 GTP-binding protein [Synechocystis sp.]	ORF2136 [Marchantia polymorpha]	oxygen independent coprophorphyrinogen III oxidase [Synechocystis sp.]	protein-export membrane protein (secF) [Haemophilus influenzae]	SPERMIDINE/PUTRESCINE TRANSPORT SYSTEM PERMEASE PROTEIN POTC.	DJ-1 protein [Homo sapiens]	DD-carboxypeptidase [Bacillus subtilis]	mating type a-1 protein [Neurospora crassa]	[TRAB [Plasmid pPD1]	GLUTAMYL-TRNA SYNTHETASE (EC 6.1.1.17) (GLUTAMATETRNA LIGASE) (GLURS).	carboxyl-terminal protease [Synechocystis sp.]	Bts1p [Saccharomyces cerevisiae]	EC 1.1.99.5 [Mus musculus]	glycerol 3 phosphate dehydrogenase [Saccharomyces cerevisiae]	glycerol uptake facilitator [Bacillus subtilis]	ORF 4 (AA 1-198); 20 kD [Escherichia coli]	putative integral membrane protease required for high frequency lysogenization by bacteriophage lambda [Escherichia coli]	HflK [Vibrio parahaemolyticus]	stringent response-like protein [Streptococcus equisimilis]	transcription elongation factor [Escherichia coli]	basic membrane protein precursor [Treponema pallidum]	H. influenzae predicted coding region HI0594 [Haemophilus influenzae]	pantothenate metabolism flavoprotein (dfp) [Haemophilus influenzae]	ORF2 gene product [Bacillus subtilis]	hypothetical protein [Synechocystis sp.]	IIPIDle2118 50S ribosomal protein L21 [Odontella sinensis]
burgdorferi - Pu gil1652022	gil11665	<u>6</u> 6	gil1573204	spIP45169IPO TC_HAEIN	.20	gil143439	gil293954	gil1041116	spiP15189ISY (E_RHIME (gil1652577	gil1098641	gil1339938	gil763191	gil142997	gil41497	gil436158	gil507734	gil407881	gi	gil	gil1573583	gil1573978	gil49316	gil1001473	gnllPIDle2118 48
Borrelia bu 90194 gil	197862	213956	217193	231762 spl	261614	278735	325818	482759	528801	543747	589360 gil	660623	660735 gi	664159	702631 gil4	704671	705643 gil	713019	783289	804832 gil	7467	.51786 gil		19767	91806 gnl
91198	198041	214639	218116	230965	262171	279964	326012	484000	527314	542317	590442	660784	662231	664938	702035	705645	706431	715040	780572	803786	8945	50587	67740	78979	92123
77	509	227	232	247	272	290	333	208	553	569	620	701	702	704	746	748	749	756	825	853	4	42	57	64	80
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	34	44	39	42	46	41	32	36	28	45	40	37	36	40	41	29	35	45	45	48	40	42	37	37		29
	62	62	62	62	62	62	62	62	62	62	62	62	62	62	62	62	61	61	61	61	61	61	61	61		61
Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins	hypothetical protein [Synechocystis sp.]	predicted 12.5Kd protein [Mycobacteriophage 15]	ribose 5-phosphate isomerase [Synechocystis sp.]	similar to APE1/LAP4, vacuolar aminopeptidase [Saccharomyces cerevisiae]	cysteinyl-tRNA synthetase [Bacillus subtilis]	similar to proofreading 3'-5' exonuclease and polymerase [Treponema pallidum]	putative orfW gene product [Clostridium acetobutylicum]	spoO193 gene product [Bacillus subtilis]	cheB peptide [Escherichia coli]	phosphomannose isomerase [Escherichia coli]	single-stranded-DNA-specific exonuclease (rec1) [Haemophilus influenzae]	unknown [Helicobacter pylori]	protoporphyrinogen oxidase (hemK) [Haemophilus influenzae]	hypothetical protein [Synechocystis sp.]	phosphatidate cytidylyltransferase [Synechocystis sp.]	collagenase [Clostridium perfringens]	tRNA guanine transglycosylase [Zymomonas mobilis]	adenine phosphoribosyltransferase form 1 [Triticum aestivum]	TagE [Vibrio cholerae]	hypothetical protein [Synechocystis sp.]	endospore forming protein [Bacillus subtilis]	gene not found in Erwinia uredovora crt gene cluster; ORF6 [Erwinia herbicola]	210668 splP37214IER GTP-BINDING PROTEIN ERA HOMOLOG. A. STRMU	possible N-terminal signal sequence; mature protein may be	membrane-anchored and start at Cys-17. 17.5% identity over 354-aa overlap with Candida pelliculosa beta-glucosidase.; putative IBacillus subtilis1	ORFveg 110 [Dictyostelium discoideum]
burgdorferi - P	gil16526/9	gil15893	gil1001678	gil529118	gil289284	352714 gil1633576	gil312380	gil40031	gil145524	$\dot{\mathbf{g}}$	93.	gil1477770	gil1574130	gil1652444	gil1652668	gil440851	gil498141	gil726305	gil460955	gil1001126	gi	gil148409	spIP37214IEF A_STRMU	gil438455		gil1513240
Borrelia	106/93	107883	213969	253175	287274 gil	352714	422495 gil	459582	484494	492322	644598	655063	714979 gi	719198	793891	862737	54726 gil	92174	106557		153051	184227	210668	264062		265581
1071201	10/458	107464	213238	251889	288749	349982	423190	458740	485147	491201	646727	655800	715668	718374	792941	862498	55889	92710	106820	111699	154445	185315	209790	262392		265982
101	IOI	102	226	266	299	357	443	489	511	518	685	695	758	762	837	917	46	81	100	109	157	193	223	273		277
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611 43			61 38	61 44	61 47	61 31	61 44	61 29	61 32	61 38	61 26	61 41	61 50	61 45	61 43	61 36	60 45	60 45	60 49	60 33	60 37	60 37	60 40	60 40	60 57	60 26	77
Pedorferi - Putative coding regions of novel proteins similar to know proteins 1354776 IMCP-1 (Trengens pallidum)				protein L17 [Escherichia coli]	ORF [Sulfolobus shibatae]	ribosomal protein L30 - Bacillus stearothermophilus	ribosomal protein L18 - Bacillus stearothermophilus	prolipoprotein signal peptidase [Staphylococcus aureus]	ssing protease [Saccharomyces cerevisiae]		TpN38(b) [Treponema pallidum]	dnaK homologue [Borrelia burgdorferi]	chocystis sp.]	YqgP [Bacillus subtilis]	hydrolase (GB:Z33006_1) [Haemophilus influenzae]	YqfM [Bacillus subtilis]	T24A11.1 [Caenorhabditis elegans]	YqgR [Bacillus subtilis]	s jannaschii]	hypothetical protein (SP:P32720) [Mycoplasma genitalium]) [Mycoplasma capricolum]	OrfH [Borrelia burgdorferi]		UDP-N-acetylglucosamine 1-carboxyvinyltransferase [Bacillus 6 subtilis]	tein diacylglyceryl transferase (lgt) [Haemophilus	us francisci]	
rferi - Putative coding	1737743 RecG [Tr			ribosomal riposomal	98	27IR	02IR			2				1303863 YqgP [Ba	573586 hydrolase		19		591369 cytidylate			372995 OrfH [Bor		PID e2768 UDP-N-ac subtilis	573923 prolipoprolinfluenzae	П	
Borrelia bu	313338 mil	1300010	372392 gi	4 401479 gil	7 404444 gil	405616 pir 5B	406435 pir 5B		441042 gil	581547 gil	585476 gil		635469 gil	(lig /59869	846688 gil 1	883282 gil	10627 gn 90	30475 gil l	44267 gil1	192053 gil	339440	362233 gil1	401872 gil	418793 gn 30	539698 gill	559655 gill	
701035				408 401874		415 405927	17 406848	41 421784	57 440722		15 584397	73 632123	675 634207	13 699438	97 847575	38 882836	7 10415	23 31428	35 44812	192994	17 341167	361817	9 402924	38 420142	540696	37 559368	000173
02 1	1 301	7C T	1 38	1 4(1 413	1 4	1 417	I 44	1 467	1 613	1 615	1 673	. 1 67	1 743	1 897	1 938	T	1 2	1 3	1 198	1 347	1 369	1 409	1 438	1 566	1 587	11

98/	589	43)	I	PCT/U	JS9	8/127	64
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47	34	33	56	38	36	38	Š	34	30	4	38	40	33		42	41	38	98	34	35	20		40	35	39	38	38	34
09	09	09	09	09	09	09		3	9	9	59	29	65		59	59	59	59	59	29	20	77	59	29	59	59	59	59
- Putative coding regions of novel proteins similar to know proteins O hypothetical protein [Synechocystis sp.]	elongation factor P [Synechococcus PCC7942]	hypothetical [Haemophilus influenzae]	mxaC gene product [Methylobacterium extorquens]	cytidylate kinase [Mycoplasma genitalium]	hypothetical [Haemophilus influenzae]	2550 hypothetical protein [Bacillus subtilis]		NifS protein. [Escherichia coli]	unknown [Schistosoma mansoni]	[type-I signal peptidase SpsB [Staphylococcus aureus]	ORF6 gene product [Bacillus subtilis]	DNA polymerase III subunit [Bacillus subtilis]	dipeptide transport system permease protein (dppB) [Haemophilus	influenzae]	phosphoglucose isomerase (AA 1-549) [Escherichia coli]	hypothetical [Haemophilus influenzae]	exodeoxyribonuclease V (recB) [Haemophilus influenzae]	rep helicase, single-stranded DNA-dependent ATPase (rep) homolog - Haemophilus influenzae (strain Rd KW20)	ORF_f560 [Escherichia coli]	Similar to arginyl-tRNA synthetase (E. coli) [Saccharomyces	CCIEVISIAE	allemate gene name yiod (eschencina com)	ORF for L15 ribosomal protein [Bacillus subtilis]	sigma factor (ntrA) (AA 1-502) [Azotobacter vinelandii]		unknown [Bacillus subtilis]	regulatory components of sensory transduction system [Synechocystis sp.]	proton glutamate symport protein [Bacillus caldotenax]
Borrella burgdorteri - Pt 690076 gil 1001260	691659 gill 399829	706626 gil1573060	gill 164996	gil1046033	813773 gill 574569	816105 gnllPIDle255	93	gil1742766	874110 gil1002666	882861 gil1595810	63234 gil580902	103802 gil467409	119914 gil1574678		141174 gil42377	gil1573129	245713 gil1574781	276257 pirID64084ID 64084	gil882504	296707 gil487937	יואררייבי	324004 g11400733	405179 gil216338	gil39269	461411 pirlA301911A 30191	463752 gil467425	481016 gil1651878	496395 gil143002
80rrelia 690076	69169	706626	735635	786567	813773	816105		829943	874110	882861	63234	103802	119914	-	141174	187659	245713	276257	281525	296707	172100	324304	405179	440759 gil39269	461411	463752	481016	496395
126069	691078	707879	734589	į .		813727		831250	872578	882211	63629	102744	118925		139567	186577	242174	278281	280005	294923	10000	972004	405646	439470	462064	462955	480078	16767
736	738	750	784	829	862	863		878	929	937	54	96	120		140	195	259	288	291	306		332	414	465	492	495	503	503
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			Borreli	burgdorferi - Puta	Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins		
	941	885060	886019	gil1685110	tetrahydrofolate dehydrogenase/cyclohydrolase [Streptococcus thermophilus]	59	36
	40	50348	48951	1574003	pantothenate permease (panF) [Haemophilus influenzae]	58	38
Ī	76	90160	89534 gil	303791	YgeJ [Bacillus subtilis]	28	32
	116	115845	115654 gnll	PIDle2758	T06E6.f [Caenorhabditis elegans]	28	37
	179	173515	173009	573163	hypothetical [Haemophilus influenzae]	58	37
	197	191904	189634	gil1066850	putative [Rhodobacter capsulatus]	28	37
	229	215111	214563	gil1573441	oxygen-independent coproporphyrinogen III oxidase (hemN) [Haemophilus influenzae]	28	34
	257	238952	241873	gil1041785	rhoptry protein [Plasmodium yoelii]	28	30
	440	421010	420792		(AE000047) Mycoplasma pneumoniae, MG246 homolog, from M. genitalium [Mycoplasma pneumoniae]	28	37
	557	533653	534750	534750 gil974332	NAD(P)H-dependent dihydroxyacetone-phosphate reductase [Bacillus subtilis]	58	41
	989	557259	559370	gil153062	helicase [Staphylococcus aureus]	28	41
	623	591542	592435	<u>.</u>	hypothetical protein [Synechocystis sp.]	28	35
	728	683208	684104	gil790935	fliG [Treponema denticola]	58	31
	961	750629	749508 gil	1574412	alanine racemase, biosynthetic (alr) [Haemophilus influenzae]	58	29
	823	778475	778723	1209836	minus strand repeat motif-containing gene [Borrelia burgdorferi]	58	22
	830	786540	788225	gil1574150	ribosomal protein S1 (rpS1) [Haemophilus influenzae]	58	34
	842	796255	796019	gnIIPIDle2434 74	ORF YGR089w [Saccharomyces cerevisiae]	58	35
	883	834332	834520	gil1575792	low Mr GTP-binding protein Rab32 [Homo sapiens]	28	43
	905	853953	854435	gil1303823	YqfG [Bacillus subtilis]	28	34
	919	863594	862875	gil1256625	putative [Bacillus subtilis]	28	34
	921	865297	864725	gil1054584	putative protein highly homologous to E. coli RNase HII [Magnetospirillum sp.]	58	42
	196	189636	187702	4	tlpC gene product [Bacillus subtilis]	57	32
Ī	797	249142	248192	gil46605	lacC polypeptide (AA 1-310) [Staphylococcus aureus]	57	41
	311	300776	301660	gil467431	high level kasgamycin resistance [Bacillus subtilis]	57	35
	365	358725	358495	gil396943	early protein [Human papillomavirus type 19]	57	38
	386	378249	378025	gil45986	NAD synthetase [Rhodobacter capsulatus]	57	32

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low proteins 437 [Methanococcus													1		l	56	56	56	95	56	56	56	56		1
Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins 394247 gil 1592085 M. jannaschii predicted coding region MJ 1437 [Methanococcus	jannaschii]	GTP-binding protein [Treponema pallidum]	Ribosomal Protein L10 [Bacillus subtilis]	YqgH [Bacillus subtilis]	ORF2136 [Marchantia polymorpha]	acriflavine resistance protein (acrB) [Haemophilus influenzae]	histidyl-tRNA synthetase [Methanococcus jannaschii]	elongation factor Ts [Chlamydia trachomatis]	hypothetical [Haemophilus influenzae]	50S ribosomal subunit protein L9 [Escherichia coli]	replicative DNA helicase [Synechocystis sp.]	acetyl coenzyme A acetyltransferase (thiolase) (fadA) homolog - Haemophilus influenzae (strain Rd KW20)	M. jannaschii predicted coding region MJ0798 [Methanococcus jannaschii]	ORF4 [Bacillus subtilis]	phospholipase C (EC 3.1.4.3) precursor - Clostridium bifermentans	exonuclease SbcD [Escherichia coli]	probable com101A gene [Haemophilus influenzae]	large tegument protein [Human herpesvirus 7]	ORF YPL216w [Saccharomyces cerevisiae]	NADH oxidase [Serpulina hyodysenteriae]	M. jannaschii predicted coding region MJ0240 [Methanococcus jannaschii]	aminodeoxychorismate lyase (pabC) [Haemophilus influenzae]	xylose repressor [Bacillus subtilis]	red alga1 chloroplast [Plasmodium falciparum]	UDP-N-acetylmuramoylalanine-D-glutamate ligase (murD) [Haemophilus influenzae]
Borrelia burgdorferi - Putati 394247 gil 1592085 N	e j	gil1732241	gil786163	gil1303855	gil11665	gil1573914	gil1591660	gil1518661	573941	gil537044	001271	IA I	806952 gil1499620 M	867809 gil1237015 O	6 pirlB30565lB 30565	gil1657594		39633	*IDle2469	2030	99018	73431	gil143841	gnIIPIDle2202 40	306992 gil1574691 UI
394690		397512	504504	689992	745857	768735	776835	790907	792328	98066 <i>L</i>	200668	802510	805240	865347	17611	35530	68915	91821	113768	142606	148561	165431	176655	301170	308362
1 399		1 402	1 533	1 735	1 794	1 814	1 821	1 834	1 836	1 848	1 849	1 851	1 855	1 922	1 12	1 26	1 59	1 79	1 112	1 147	1 153	1 169	1 183	1 312	1 317

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	448	426477	Borrelia 426133	burgdorferi - Pu	Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins 26133 gil467410 Junknown [Bacillus subtilis]	98	28
	456	437678	434457	oil142521	deoxyrihodinyrimidine nhotolyase [Bacillus subtilis]	25	
	500	070764	10101	1777119	decolationally minimum photoly asc [Dacinius submits]	200	1
-	94	4381/8	43/312 8118	g11882453	ORF_1286; alternate name yggB; or14 of X14436 [Escherichia colii]	26	31
	160	441300	1/3/120 ail 1	n:11/8216	NoII ontinorter protein [Entercoccons hine]	75	22
	407	441302	447470	100	INAIT-AIRIDUILEI DIOLEIN [EMICIOCOCCUS IIIIAE]	20	37
1	809	574772	574951	gill	NADH dehydrogenase subunit 2 [Paramecium aurelia]	99	37
1	669	659498	660055	gill	OrfH [Borrelia burgdorferi]	99	24
-	757	713509	713712	gil861327	F31D5.5 gene product [Caenorhabditis elegans]	99	40
-	791	741305	742837	gil1651873	4-alpha-glucanotransferase [Synechocystis sp.]	56	43
	822	779478	778291	gil1500309	M. jannaschii predicted coding region MJ1428 [Methanococcus	56	28
					Jannascnii		
-	196	907556	908932	gil1749528	similar to Saccharomyces cerevisiae probable UTP-glucose-1-	99	37
					phosphate uridylyltransferase, SWISS-PROT Accession Number D32861 [Schizosacharomyces nombel	************	
					r 32001 [Scilledsaccidalulilyces pullide]		
1	39	48953		gil1045895	[hypothetical protein (SP:P23851) [Mycoplasma genitalium]	55	41
T	131	132989	131967	gil1574007	nitrogen fixation nifR3 protein (nifR3) (PIR:S49971)	55	39
					[Haemophilus influenzae]		
1	152	148506	147148	gil1653100	Na+ -ATPase subunit J [Synechocystis sp.]	55	31
1	359	352690	353313 gill.	gi11213334	OrfX; hypothetical 22.5 KD protein downstream of type IV	55	33
					prepilin leader peptidase gene; Method: conceptual translation		
					supplied by author [Vibrio vulnificus]		
1	361	355510	354140	gil882698	L-fuculose kinase [Escherichia coli]	55	44
1	515	488398	487652	gil397486	endonuclease G [Bos taurus]	55	33
1	551	526427	525285	gil558266	orf gene product [Wolinella succinogenes]	55	30
I	270	543745	544482 gil]	gil1303811	YqeU [Bacillus subtilis]	55	33
1	279	551201	551494 ₈	gil290487	50S ribosomal subunit protein L28 [Escherichia coli]	55	37
1	584	555359	556063 ₈	gil1592301	M. jannaschii predicted coding region MJ0687 [Methanococcus	55	32
					jannaschii]		
	902	665310	965936		deoxyguanosine kinase/deoxyadenosine kinase(I) subunit [Lactobacillus acidophilus]	55	38
	771	722876	723538	gil1736440	O-sialoglycoprotein endopeptidase (EC 3.4.24.57) (Glycoprotease). [Escherichia coli]	55	39
-	786	736537	737187	gil1589778	SPINDLY [Arabidopsis thaliana]	55	34
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823341 823790 gil1590959 ĀTP synthase, subunit K [Methanococcus jannaschii] 55 847660 84942 gil1517942 aminopeptidase P [Sus scrofa] 55 867811 868236 gil114260 POMI [Plasmodium chabaudi chabaudi] 55 870915 870005 870005 870005 870005 870916 8700390 gil312694 ARS-binding factor 1 [Kluyveromyces maxianus] 55 44068 74679 gil41736 Orf635 gene product [Euglena gracilis] 54 184282 182904 gil151259 MMG-CoA reductase (EC. 1.1.188) [Pseudomonas mevalonii] 54 194105 192951 gil1045800 ribose transport system permease protein [Mycoplasma genitalium] 54 210749 21230 gil1591243 M. jamaschii] predicted coding region MJ0339 [Methanococcus 54 237491 238954 gil169164580 m. jamaschiii predicted coding region MJ0263 [Methanococcus 54 237491 238046 M. jamaschiii predicted coding region MJ0263 [Methanococcus 54 37706 379909 gil1909258 D.Dcarboxypeptidase [Eucroccus faccalis] 54 37706 37726 gil1439647 M. jamaschiii predicted coding reg	11 810	765243	Borrelia burgo 766130 gil98	dorferi - Puta 14805	rigdorferi - Putative coding regions of novel proteins similar to know proteins 1984805 [Blycine betaine-binding protein precursor [Bacillus subtilis]	55	35
847660 849462 gill 15 17942 aminopopidase P (Sus scrotal) 55 887811 88623 gill 15 17942 aminopopidase P (Sus scrotal) 55 870905 870079 gill 14566 POMJI (Plasmodium chabadi chabaudi) 55 904091 903900 gill 23839 CheR (Rhizobium melitoti) 54 44068 43124 gill 48660 delta-2-isopentenyl pyrophosphate transferase [Escherichia coli] 54 44068 43124 gill 48860 delta-2-isopentenyl pyrophosphate transferase [Escherichia coli] 54 79094 74709gill 47757 Grids 5 gene producte [Euglena gardlis] 54 194105 192909 gill 501243 M. jannaschii predicted coding region MJ0539 [Methanococcus 34 54 210749 212320 gill 591243 M. jannaschii predicted coding region MJ056 [Methanococcus 34 54 237491 238954 gill 791242 M. jannaschii predicted coding region MJ056 [Methanococcus 34 54 311333 312133 gill 209228 Dcaboxypeptidase [Enerococcus faecalis] 54 37096 377096 377096 377096 377096 377096 377096 377096 377096 377	1 871	823341	. 20	656069	ATP synthase, subunit K [Methanococcus jannaschii]	55	34
867811 868236 gill 142660 POM1 [Plasmodium chabaudi chabaudi] 55 870905 87005 970059 Gil12694 ARS-binding factor 1 [Kluyveronyces marxianus] 55 904091 903900 gil12694 ARS-binding factor 1 [Kluyveronyces marxianus] 54 40608 43124 gil46860 delta-2-isopentenyl pyrophosphate transferase [Escherichia colij] 54 184282 182969 gil15129 HMG-CoA reductase (EC 1.1.88) [Pseudomonas mevalonii] 54 194105 192951 gil164860 delta-2-isopentenyl pyrophosphate transferase [Escherichia colij] 54 194105 192951 gil164860 delta-2-isopentenyl pyrophosphate transportenyl pyrophosphatese [Entercococcus political pyrophatese [Entercococcus political pyrophatese] 54 24586 247542 gil1874782 exodeoxyribonuclease V (recD) [Haemophilus influenzae] 54 577096 579909 gil1499043 M. jannaschii predicted coding region MI026 [Methanococcus pyropenial pyropenia	1 898	847660	. <u>5</u> 2	17942	aminopeptidase P [Sus scrofa]	55	46
870905 870039 gil534839 CheR (Rhizobium melitot) 904091 903900 gil512860 delta-2-isopenteuryl pyropkente transferase [Escherichia coli] 55 904091 903900 gil312860 delta-2-isopenteuryl pyropkoshate transferase [Escherichia coli] 54 79004 74679 gil415736 Orf635 gene product [Euglena gracilis] 54 79034 74679 gil415736 Orf635 gene product [Euglena gracilis] 54 194105 192951 gil15739 HMG-CoA reductase (EC 1.1.1.88) [Pseudomonas mevalonil] 54 194105 192320 gil1591243 M. jannaschii predicted coding region MJ0339 [Methanococcus familiana) 54 245698 247542 gil1574782 exodeoxyribonuclease V (recD) [Haemophilus influenze] 54 247698 247542 gil1574782 exodeoxyribonuclease V (recD) [Haemophilus virilis] 54 377096 579909 gil1499043 M. jannaschii predicted coding region MJ0263 [Methanococcus familianaschii] 54 797932 79999 gil290216 [bride of sevenless] gene product [Drosophila virilis] 54 884888 893912 gil173804 dosage-dependent dnak suppressor protein [Escherichia coli] 54 96019 <td< td=""><td>1 924</td><td>867811</td><td><u></u></td><td>42660</td><td>_</td><td>55</td><td>41</td></td<>	1 924	867811	<u></u>	42660	_	55	41
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44068 43124 gil 146860 delta-2-isopentenyl pyrophosphate transferase [Escherichia coli] 54 79094 74679 gil 14536 Orf655 gene product [Euglena gracilis] 54 179094 74679 gil 151259 HMO-Co-Ar ceductase (EC. I.1.1.88) [Pseudomonas mevalonii] 54 194105 192951 gil 1945800 ribose transport system permease protein [Mycoplasma genitalium] 54 210749 212320 gil 1591243 M. Jannaschii predicted coding region MJ0539 [Methanococcus and protein [Mycoplasma genitalium] 54 237491 238954 gnllPIDE-2450 unknown [Mycobacterium tuberculosis] 54 245698 247542 gil 1574782 exodeoxyribonuclease V (recD) [Haemophilus influenzae] 54 245698 247542 gil 1499043 M. Jannaschii predicted coding region MJ0263 [Methanococcus 54 54 577096 579909 gil 1499043 M. Jannaschii predicted coding region MJ078 [Methanococcus 54 54 720685 719999 gil 1499043 M. Jannaschii predicted coding region MJ0798 [Methanococcus 54 54 739607 739996 gil 1495043 M. Jannaschii predicted coding region MJ0798 [Methanococcus 54 54 79931 768386 M. Jannaschii predicted coding region MJ0798 [Methanococc	1 964	904091	903900 gil31	2694	ARS-binding factor 1 [Kluyveromyces marxianus]	55	50
79094 74679 gil415736 Orf635 gene product [Euglena gracilis] 184282 182969 gil415739 HMG-CoA reductase (EC I.1.1.88) [Pseudomonas mevalonii] 54 184282 182969 gil151239 HMG-CoA reductase (EC I.1.1.88) [Pseudomonas mevalonii] 54 194105 192931 gil1045800 ribose transport system permease protein [Mycoplasma genitalium] 54 210749 212320 gil1591243 M. jannaschii predicted coding region MJ0539 [Methanococcus 54 245698 247542 gil1591245 unknown [Mycobacterium tuberculosis] 247542 gil1309542 D.D-carboxypeptidase [Enterococcus faecalis] 54 577096 579909 gil1499043 M. jannaschii predicted coding region MJ0263 [Methanococcus 54 577096 579909 gil1499043 M. jannaschii predicted coding region MJ0263 [Methanococcus 54 577096 247542 gil1499043 M. jannaschii predicted coding region MJ0708 [Methanococcus 54 577096 579909 gil1499043 M. jannaschii predicted coding region MJ0798 [Methanococcus 54 579909 gil1499043 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 54 579909 gil1499650 Ingeallar P-ring protein [Pseudomonas putida] 55 570909 gil1499650 Ingeallar P-ring protein [Pseudomonas putida] 55 57090 gil1499650 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 5207426 gil1499650 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 5207426 gil1499650 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 5207426 gil1499650 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 5207102 gil1499620 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 57009 gil1015945 methyl accepting chemotaxis homolog [Treponena denticola] 55 57009 gil1015945 methyl accepting chemotaxis homolog [Treponena denticola] 55 57009 gil1015945 methyl accepting chemotaxis homolog [Treponena denticola] 57000 gil1015945 methyl accepting chemotaxis homolog [Treponena denticola] 57000 gil10160000000000000000000000000000000000	1 33	44068	43124 gil14	0989	delta-2-isopentenyl pyrophosphate transferase [Escherichia coli]	54	31
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194105 192951 gil1045800 ribose transport system permease protein [Mycoplasma genitalium] 54 210749 212320 gil1591243 M. jannaschii predicted coding region MJ0539 [Methanococcus jannaschii] 54 237491 238954 gnllPIDle2450 unknown [Mycobacterium tuberculosis] 54 245698 247542 gil1574782 exodeoxyribonuclease V (recD) [Haemophilus influenzae] 54 245698 247542 gil1574782 exodeoxyribonuclease V (recD) [Haemophilus influenzae] 54 311333 312133 gil1209528 D.D-carboxypeptidase [Enterococcus faecalis] 54 377096 379909 gil290216 Ibride of sevenless] gene product [Drosophila virilis] 54 720685 719999 gil290216 Ibride of sevenless] gene product [Drosophila virilis] 54 720687 739906 gil473604 'dosage-dependent dnak suppressor protein' [Escherichia coli] 54 797932 798360 Janaschiils subtilis} 54 96019 97032 gil405560 Ilagellar P-ring protein [Pseudomonas putida] 53 98331 199215 gil1303842 Yqft [Bacillus subtilis] 53 159533 158562 gil1499650 M. jannaschii predicte	1 192	184282	182969 gill	1259	(EC 1	54	35
210749 212320 gil1591243 M. jannaschii predicted coding region MJ0539 [Methanococcus jannaschii] 54 237491 238954 gillPDle2450 unknown [Mycobacterium tuberculosis] 54 245698 247542 gill274782 exodeoxyribonuclease V (recD) [Haemophilus influenzae] 54 311333 312133 gill209228 D.D-carboxypeptidase [Enterococcus faecalis] 54 577096 579909 gill499043 M. jannaschii predicted coding region MJ0263 [Methanococcus jannaschii] 54 720685 719999 gil290216 [bride of sevenless] gene product [Drosophila virilis] 54 739607 739996 gil473804 dosage-dependent dnaK suppressor protein [Escherichia coli] 54 797932 798366 gil473804 dosage-dependent Beculomonas putida] 54 86019 97032 gil405767 riposomal protein [Pseudomonas putida] 54 96019 97032 gil405550 flagellar P-ring protein [Pseudomonas putida] 53 98331 99115478 No definition line found [Escherichia coli] 53 159533 158562 gil499620 M. jannaschii predicted coding region MJ0798 [Methanococcus standarding line found line found line found line giland line found line found line found line found line found	1 200	194105	192951 gill(008540	ribose transport system permease protein [Mycoplasma genitalium]	54	29
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245698 247542 gill 574782 exodeoxyribonuclease V (recD) [Haemophilus influenzae] 54 311333 312133 gill 209528 D.D-carboxypeptidase [Enterococcus faecalis] 54 577096 579909 gill 499043 M. jannaschii predicted coding region MJ0263 [Methanococcus jannaschii] 54 720685 719999 gill 200216 [bride of sevenless] gene product [Drosophila virilis] 54 739607 739996 gill 47804 dosage-dependent dnaK suppressor protein [Escherichia coli] 54 797932 798366 gill 045767 ribosomal protein [Se [Mycoplasma genitalium] 54 894898 893912 gill 303842 YqU [Bacillus subtilis] 53 96019 97032 gill 409550 flagellar P-ring protein [Pseudomonas putida] 53 159533 158562 gill 1499620 M. jannaschii predicted coding region MJ0798 [Methanococcus familar to Saccharomyces cerevisiae unknown, EMBL Accession 53 266053 267426 gill 749686 similar to Saccharomyces cerevisiae unknown, EMBL Accession 53 292150 294309 gill 1015945 methyl accepting chemotaxis homolo	1 256	237491	gn 24	*IDIe2450	unknown [Mycobacterium tuberculosis]	54	34
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96019 97032 gil405550 flagellar P-ring protein [Pseudomonas putida] 53 98331 99215 gil912478 No definition line found [Escherichia coli] 53 159533 158562 gil1499620 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 234276 232861 gil1303989 YqkI [Bacillus subtilis] 53 266053 267426 gil1749686 similar to Saccharomyces cerevisiae unknown, EMBL Accession 53 Number Z68194 [Schizosaccharomyces pombe] Number Z68194 [Schizosaccharomyces pombe] 53 292150 294309 gil1015945 methyl accepting chemotaxis homolog [Treponema denticola] 53 358298 357702 gil1499620 M. jannaschii predicted coding region MJ0798 [Methanococcus 53	1 951	894898	893912 gill 3	03842	YqfU [Bacillus subtilis]	54	28
98331 99215 gil912478 No definition line found [Escherichia coli] 53 15953 158562 gil1499620 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 234276 232861 gil1303989 YqkI [Bacillus subtilis] 53 266053 267426 gil1749686 similar to Saccharomyces cerevisiae unknown, EMBL Accession 53 Number Z68194 [Schizosaccharomyces pombe] Number Z68194 [Schizosaccharomyces pombe] 53 358298 357702 gil1499620 M. jannaschii predicted coding region MJ0798 [Methanococcus 53 jannaschii] jannaschiil	1 86	96019	97032 gil40	5550	flagellar P-ring protein [Pseudomonas putida]	53	40
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234276232861 gil1303989YqkI [Bacillus subtilis]53266053267426 gil1749686similar to Saccharomyces cerevisiae unknown, EMBL Accession53Number Z68194 [Schizosaccharomyces pombe]53292150294309 gil1015945methyl accepting chemotaxis homolog [Treponema denticola]53358298357702 gil1499620M. jannaschii predicted coding region MJ0798 [Methanococcus53	1 164	159533	158562 gil	99620	M. jannaschii predicted coding region MJ0798 [Methanococcus annaschii]	53	39
266053267426 gil1749686similar to Saccharomyces cerevisiae unknown, EMBL Accession53292150294309 gil1015945methyl accepting chemotaxis homolog [Treponema denticola]53358298357702 gil1499620M. jannaschii predicted coding region MJ0798 [Methanococcus]53	1 250	234276	232861 gil	68650	YqkI [Bacillus subtilis]	53	28
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	1 364	358298	357702 gil 14		M. jannaschii predicted coding region MJ0798 [Methanococcus annaschii]	53	41

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28 33 32 35 30 26	26 34 37 24	34 25 28 28 32 47 47 40	35 29 20 27 27 29 29 29 29 29
53 53 53	22 22 22	\$25 \$2 \$2 \$25 \$2 \$25	\$25 \$25 \$25 \$25 \$25 \$25 \$25 \$25 \$25 \$25
Putative coding regions of novel proteins similar to know proteins orf 06111 gene product [Saccharomyces cerevisiae] YIXH [Borrelia burgdorferi] cell division protein J [Methanococcus jannaschii] GlcNAc 6-P deacetylase [Vibrio furnissii] YqhZ [Bacillus subtilis] H. influenzae predicted coding region HI1555 [Haemophilus	influenzae] hypothetical protein [Synechocystis sp.] P35 gene product (AA 1 - 314) [Escherichia coli] repeat organellar protein [Plasmodium chabaudi] colicin V production protein (pur regulon) (cvpA) [Haemophilus influenzae]	secA gene product [Antithamnion sp.] hypothetical protein [Synechocystis sp.] trigger factor [Bacillus subtilis] glutamic acid-rich protein [Plasmodium falciparum] 24K membrane protein [Pseudomonas aeruginosa] phnP protein [Escherichia coli] unknown [Bacillus subtilis] hypothetical protein (GP:X91006_2) [Methanococcus jannaschii] (AE000047) Mycoplasma pneumoniae, MG246 homolog, from	M. genitalium [Mycoplasma pneumoniae] aspartyl-tRNA synthetase (aspS) [Haemophilus influenzae] fibronectin/fibrinogen-binding protein [Streptococcus pyogenes] dihydroorotate dehydrogenase [Plasmodium falciparum] S2 gene product [Borrelia burgdorferi] SpoVD [Bacillus subtilis] ATP synthase, subunit D [Methanococcus jannaschii] repeat organellar protein [Plasmodium chabaudi] putative [Bacillus subtilis]
Borrelia burgdorferi - P 486888 gil940842 540684 gil1165254 591032 gil1592021 758537 gil1732203 805298 gil1303915 834944 gil1574399	56944 gil1652686 62383 gil42219 65665 gil1151158 102746 gil1574136	116879 gil288998 208446 gil1652602 272764 gnllPIDle255 28 346532 gil160299 361800 gil216861 367695 gil147213 372412 gil467459 416768 gil1591425	443798 gil1573287 553802 gil496254 715610 gil397703 750674 gil1063419 774852 gil580936 821516 gil1592298 838106 gil1151158 862110 gil1256625
486253 541832 590418 759748 804825 835705	58236 63264 66168 102255	115800 208898 274152 344946 361087 368462 373209 418141 420801	443436 555235 715852 751384 776768 820887 839581 862856 83112
514 567 621 805 884 884	53 56 95	220 220 285 362 368 376 381 437 439	583 759 797 820 869 888 888 916

	1	1				1 29	51 26			50 25	50 35	50 29	50 32			50 30		50 32			50 30		48 21				70 21
	[5	5	5	5	5	[2	5	5	5	5	5	5	2	5	5	5	5	5	5	5	5	4	4	4	4	4	V
ative coding regions of novel proteins similar to know proteins	20844 orf4 [Bacillus subtilis]	3-hydroxy-3-methylglutaryl-CoA synthase [Gallus gallus]	ORF2 [Bacillus subtilis]	protein antigen LmSTI1 [Leishmania major]	chromate resistance protein A [Methanococcus jannaschii]	PIDIe2390 AMP-binding protein [Brassica napus]	a negative regulator of pho regulon [Pseudomonas aeruginosa]	ORF2 [Salmonella typhimurium]	phospho-N-acetylmuramoyl-pentapeptide- transferase [Bacillus subtilis]	RING-finger protein [Helicoverpa armigera nucleopolyhedrovirus	PgsA [Bacillus subtilis]	YqfV [Bacillus subtilis]	PIDle2767 unknown [Mycobacterium tuberculosis]	peptidase D [Escherichia coli]	ComE [Synechocystis sp.]	PIDIe2202 frameshift [Plasmodium falciparum]	beta-galactosidase [Thermoanaerobacterium thermosulfurigenes]	B.subtilis genes rpmH, rnpA, 50kd, gidA and gidB [Bacillus subtilis]	CG Site No. 29739 [Escherichia coli]	T03G11.2 gene product [Caenorhabditis elegans]	murE gene product [Bacillus subtilis]	involucrin [Saguinus oedipus]	ORF2 [Salmonella typhimurium]	yejE [Escherichia coli]	putative [Bacillus subtilis]	FemA [Staphylococcus simulans]	Lemosthatian I manatoin [Cumanhangration and
Borrelia burgdorferi - Put	146360 gil520844	185275 gil211931	289676 gil142833	gill	439497 gil1591434		869955 gil633996	54062 gil505363	gil3	117096 gil1762996	150506 gil893358	224744 gil1303843	,	274710 gil147140		341581 gnilPIDle2202 45	434509 gil144839	447948 gil580905	641039 gil882579	690400 gil1086864	709662 gil40162	354157 gil343314	53216 gil505363	120774 gil405908	156653 gil143213	gill	11001170
	147190	186516	288759	362209	438943	<u> </u>	869257	54760	101155	118397	151159	224187	L	276164	299525	342477	435120	448691	640194	690152	708130	353288	54046	119896	157504	305940	10/150
	150	194	300	371	464	819	976	45	94	118	155	239	274	287	310	349	457	479	089	737	752	360	44	122	161	316	
	F	-	F		F		-	F		T	-	F			-	-	F		-	-	-	-	I		F	=	ŧ

9 1	628 596267	Borrelia burgdorferi - Puta 596566 gill 56218	dorferi - Putative coding regions of novel proteins similar to know proteins 56218 putative [Caenorhabditis elegans]	48	32
1 -		654452 gill 574476	dedA protein (dedA) [Haemophilus influenzae]	48	22
1 7	731 686392	686129 gil915207	gastric mucin [Sus scrofa]	48	27
1 8		843476 gnll	PIDIe2202 frameshift [Plasmodium falciparum]	48	32
	62 74673	72196	outer membrane protein [Neisseria gonorrhoeae]	47	30
10		108780 gill		47	27
	187 1811111	180215 gil1184118	mevalonate kinase [Methanobacterium thermoautotrophicum]	47	30
1 20		196616 gill	phosphoglycolate phosphatase, chromosomal (SP:P40852) [Haemophilus influenzae]	47	21
1 2	265 251835	251098 gil1209847	repeat motif-containing gene [Borrelia burgdorferi]	47	30
1 3			uridylate kinase [Methanococcus jannaschii]	47	26
1 3	356 349581	349991 gil8	Probable essential component of the nucleoskeleton (Swiss Prot. accession number P32380) [Saccharomyces cerevisiae]	47	27
1	460559	459834 gil1592264	type I restriction enzyme [Methanococcus jannaschii]	47	34
		1	ankyrin 3 [Mus musculus]	47	53
1 5		548390 gnll	PIDIe2202 frameshift [Plasmodium falciparum]	47	27
1 72	744 701189		PIDIe1604 orfA gene product [Borrelia burgdorferi]	47	23
1 7.	755 713050	1	710765 pirlS41649IS4 DNA polymerase - Plasmodium falciparum	47	22
11 7/	761 717229	i	M. jannaschii predicted coding region MJ1428 [Methanococcus jannaschii]	47	37
~	813 767745	768737	membrane fusion protein (mtrC) homolog - Haemophilus influenzae (strain Rd KW20)	47	23
- X	824 779587	780546 gild	contains TPR domain-like repeats [Caenorhabditis elegans]	47	28
1	881 834283	833015	H. influenzae predicted coding region H11548 [Haemophilus influenzae]	47	24
1	886 837236	836199 gil887563	serine/threonine-protein kinase [Plasmodium falciparum]	47	30
1	47 57001	55880 gil1652686	hypothetical protein [Synechocystis sp.]	46	23
		156171 gill	ORF4 protein (AA 1-156) [Paramecium aurelia]	46	9

28	20	18	27	21	28		29	32	29	19	26	29			27	23	23	26	26	31	25	19
46	46	46	46	46	46		46	46	46	46	46	46			45	45	45	44	44	44	44	43
Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins 765 232829 gil 1142681 Lpp38 [Pasteurella haemolytica]	322838 gil562039	327303 gil457146	421747 gil15		545596 gil1022328	been purified and found to bind in vitro to a promoter region [Myxococcus xanthus]					852741 gil806562	909948 gil438455	membrane-anchored and start at Cys-17. 17.5% identity over 354-	aa overlap with Candida pelliculosa beta-glucosidase.; putative [Bacillus subtilis]		438949 gil687689	695295 gil 1499043 M. jannaschii predicted coding region M jannaschii]	96 98756 gil303895 ORF 8: This ORF is required for the secretion of IpaB, IpaC and IpaD IPasmid pMYSH60001	234343 gil143245			Ш
2317	323695	3290	42251	428632	545081		586903	668290	7411	843474	853463	908917			197467	438197	698657	96166	235698	668406	802490	436119
249	329	336	442	1 452	573		617	108	190	892	1 903	896			208	462	742	06	253	709	850	458

	30	27	19	24	25	26	26	20	24					28	-	19	24	24	24
	43	43	42	42	42	41	41	41	40			-		40	Ç	5	\$ 9	40	04 04
tative coding regions of novel proteins similar to know proteins	309967 pirlS17998IS1 gene COX1 intron 4 protein - yeast (Kluyveromyces marxianus var. lactis) mitochondrion (SGC2)	no score generated - score shown is bogus [Mycoplasma genitalium]	hypothetical protein (GP:X91006_2) [Methanococcus jannaschii]	hypothetical protein (SP:P32720) [Mycoplasma genitalium]	VAR1 protein [Candida glabrata]	ipa-52r gene product [Bacillus subtilis]	546581 gnllPIDle 1632 MURF2 protein (AA 1-348) [Crithidia fasciculata]	repeat organellar protein [Plasmodium chabaudi]	Similar to chromosome segregation protein Smc1p of S. cerevisiae	(GenBank accession number L00602), chromosome segregation	protein Cut3p of S. pombe (Swiss Prot. accession number	P41004), and C. elegans hypothetical proteins R13G10.1	(GenBank	neural specific DNA binding protein [Xenopus laevis]	hymothatian matain (CD.V01006 3) [Mathanasasans inmescalin	hisponiencal protein (Or.A) 1000_2) [menianococcus jannascinii]	[Mus musculus (strain C3HF/RL) ORF mRNA, complete cds.],	[Mus musculus (strain C3HF/RL) ORF mRNA, complete cds.], gene product [Mus musculus]	[Mus musculus (strain C3HF/RL) ORF mRNA, complete cds.], gene product [Mus musculus] wall-associated protein [Bacillus subtilis]
ı burgdorferi - Pu	PirlS17998IS 7998	gil1045905	gil1591425	gil1045801	gil343962	gil413976	gnIIPIDle 1637 6	gil1151158	gil1256888	•				gil1150836	oil1501425	51117/114	gil499647	429700 gil499647	gil499647
Вотелі	809967 pirlS179 7998	879701 gil	309877 gil	588672 gi	594572	101021 gil	546581	692403 gill	6796 gill					214742 gil	308377 gi		429700	429700	429700 gi
	810560	881179	311250	587863	593472	100191	545523	693458	5792	-				214440	309735		431037	431037	431037
	829	935	319	618	625	93	574	740	3					228	318		453	453	453
	T				1	-	1							Γ		,	T		

Borrelia burgdorferi - Coding regions containing know proteins

Contig ID	Orf ID Start (nt)		Stop (nt)	match acession	match gene name	percent ident	HSP nt length
	69		85018		Borrelia burgdorferi peptidyl-tRNA hydrolase- like protein (pth) gene homologue, complete cds	100	220
	70	86918	86340	gblL321441	Borrelia burgdorferi peptidyl-tRNA hydrolase- like protein (pth) gene homologue, complete cds	100	579
	71	87573	86911	gb L32144	Borrelia burgdorferi peptidyl-tRNA hydrolase- like protein (pth) gene homologue, complete cds	100	129
	124	123885		121759 gblM60802l	B.burgdorferei immunogen gene, 5' flank	66	2127
	126	127421	1	125700 emblX91965l BBATPBP	B.burgdorferi abp gene	26	284
	137	136332	139151	gblL314241	Borrelia burgdorferi (clone BbK3.11) phoA fusion protein gene, partial cds	86	248
	138		138515	gblL314241	Borrelia burgdorferi (clone BbK3.11) phoA fusion protein gene, partial cds	96	09
	165	160705	159932	gblU175911	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	100	774
	166		160703	160703 gblU175911	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	100	1902
	167	162835	162602	162602 gblU175911	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	66	232
	168	164397	162811	gblU175911	Borrelia burgdorferi primary sigma factor (rpoD) gene, complete cds	66	1216
_	210	198495		199028 gblU61498	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	86	127
	211	199527		199069 gblU614981	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	66	459
	212			199549 gblU61498	Borrelia burgdorferi CheA (cheA) gene, partial cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	66	519
	213	201455		200046 gblU61498l	Borrelia burgdorferi CheA (cheA) gene, partial	66	1410

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Borrelia burgdorferi - Coding regions containing know proteins

	cds, CheW (cheW), CheX (cheX) and CheY (cheY) genes, complete cds	neX) and Che Y	- 6	1000
-	Borrelia burgdorferi histidine kinase (cheA) gene, complete cds	kinase (cheA)	66	2601
204115 gblU62900l Borrelia bu protein (fla chemotaxis cds	Borrelia burgdorferi flagellar filament outsheath protein (flaA) gene, complete cds, and chemotaxis histidine kinase (cheA) gene, partial cds	filament outsheath cds, and heA) gene, partial	86	1059
205220 gblU62900l Borrelia bu protein (fla chemotaxis chemotaxis cds	Borrelia burgdorferi flagellar filament outsheath protein (flaA) gene, complete cds, and chemotaxis histidine kinase (cheA) gene, partial cds	filament outsheath cds, and :heA) gene, partial	86	277
220232 emblX651391 B.burgdorf BBHSP60 protein	B.burgdorferi hsp60 gene for 60kDa heat shock protein	60kDa heat shock	86	139
220594 emblX651391 B.burgdorf BBHSP60 protein	B.burgdorferi hsp60 gene for 60kDa heat shock protein	60kDa heat shock	66	1695
160	Borrelia burgdorferi groEL gene for a common antigen	ne for a common	94	416
	Borrelia burgdorferi (clone BbK fusion protein gene, partial cds	bK1.4) phoA ds	66	231
283629 emblX877251 B.burgdorf BBDNA66K D	B.burgdorferi p66 gene for 66kDa protein	kDa protein	100	1881
283683 gblM584311 Borrelia bu	Borrelia burgdorferei PCR target sequence	get sequence	93	356
	Borrelia burgdorferi methionyl tRNA synthetase (metG) gene, partial cds	I tRNA synthetase	66	345
	Borrelia burgdorferi response regulator gene, partial cds	regulator gene,	100	191
330299 gblL399651 Borrelia bu gene, com	Borrelia burgdorferi histidine kinase (cheA) gene, complete cds	kinase (cheA)	66	1361
	Borrelia burgdorferi phosphotransferase enzyme II (crr) gene, hsp90 (hptg) gene, complete cds	ransferase enzyme ne, complete cds	100	95
338830 gblU51878 Borrelia bu	Borrelia burgdorferi phosphotransferase enzyme	ransferase enzyme	66	1980

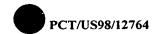
Borrelia burgdorferi - Coding regions containing know proteins

346 339458 338868 gblU51878 Borrelia burgdorferi phosphotransferase enzyme 100						II (crr) gene, hsp90 (hptg) gene, complete cds		
378955 379590 gblM968471 Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's and DnaJ protein homologue gene, complete cds s and bnaJ protein homologue gene, complete cds and thioredoxin reductase (trxB) genes, complete cds and thioredoxin reductase (trxB) gen	-	346	339458	338868	gbIU51878I	Borrelia burgdorferi phosphotransferase enzyme II (crr) gene, hsp90 (hptg) gene, complete cds	100	591
381512 emblX67646 B.burgdorferi dnak gene for heat-shock protein BBHSPRO 381512 381943 gblM97914 Borrelia burgdorferi GrpE protein homologue gene, complete cds 381507 382617 gblM96847 Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's 382656 383360 gblM96847 Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's 383005 382688 gblM96847 Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's 384408 383416 gblU82978 Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenyla	_	388	378955	379590	gbIM96847I	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	100	636
381512 381943 gblM979141 Borrelia burgdorferi DnaJ gene, complete cds 382617 gblM968471 Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's 382656 383360 gblM968471 Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's 383405 382688 gblM968471 Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's 384408 383416 gblU829781 Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl- tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheS), phenylalanyl- tRNA synthetase beta subunit (pheS) and thioredoxin reductase (trxB) genes, complete cds	-	389	379566	381521	emblX67646l BBHSPRO	B.burgdorferi dnaK gene for heat-shock protein	100	1956
381907 382617 gblM968471 Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha	-	390	381512	381943	gblM97914l	Borrelia burgdorferi DnaJ gene, complete cds	62	424
382656 383360 gblM968471 Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's protein homologue gene, complete cds's Borrelia burgdorferi GrpE protein homologue gene, and DnaJ synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase et a subunit (pheS), p	-	391	381907	382617	gbIM96847I	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	100	687
383005 382688 gblM96847l Borrelia burgdorferi GrpE protein homologue gene, and DnaJ protein homologue gene, complete cds's protein homologue gene, complete cds's synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheS), pheny	-	392	382656	383360	gbIM96847I	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	95	144
384408 383416 gblU829781 Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds thioredoxin reductase (trxB) genes, complete cds thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase alpha subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds (tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	_	393	383005	382688	gblM96847I	Borrelia burgdorferi GrpE protein homologue gene, DnaK protein homologue gene, and DnaJ protein homologue gene, complete cds's	95	144
384799 384467 gblU82978l Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheT) and tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheT) and tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds thioredoxin reductase (trxB) genes, complete cds synthetase alpha subunit (pheT) and tRNA synthetase alpha subunit (pheT) and tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds thioredoxin reductase (trxB) genes, complete cds	_	394	384408	383416	gbIU82978I	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	66	956
386169 384733 gblU82978l Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds thioredoxin phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds thioredoxin reductase (trxB) genes, complete cds		395	384799	384467	gblU82978I	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	66	292
387733 386144 gblU82978l Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	-	396	386169	384733	gblU82978I	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl- tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	66	1416
		397	387733	386144	gblU82978l	Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl- tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	66	1220

Borrelia burgdorferi - Coding regions containing know proteins

230	152	287	357	291	828	324	642
66	98	66	96	66	86	66	66
Borrelia burgdorferi phenylalanyl-tRNA synthetase alpha subunit (pheS), phenylalanyl-tRNA synthetase beta subunit (pheT) and thioredoxin reductase (trxB) genes, complete cds	B.burgdorferei promoter region DNA	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and	S3 (rpsC) gene, partial cds Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and
387727 gbiU82978I	gblM286811	410132 gblU781931	411017 gblU781931	411386 gblU781931	gbIU781931	412529 gblU78193l	412846 gblU781931
387727	407981	410132	411017	411386	411674	412529	412846
394257	408559	411019	411388	411676	412531	412852	413487
398	421	427	428	429	430	431	432
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Borrelia burgdorferi - Coding regions containing know proteins

	633	324	1212	148	171	312	180
	66	100	100	100	100	100	100
S3 (rpsC) gene, partial cds	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	Borrelia burgdorferi tuf-s10 operon: elongation factor (tuf), ribosomal proteins S10 (rpsJ), L3 (rplC), L4 (rplD), L23 (rplW), L2 (rplB), S19 (rpsS), and L22 (rplV)genes, complete cds, and S3 (rpsC) gene, partial cds	Borrelia burgdorferi elongation factor EF-Tu (tuf) gene, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA),
	413485 gbIU781931	414141 gblU781931	414503 gblL231251	450310 gblU045271	450650 gblU045271	450897 gbIU045271	451467 gbIU045271
	413485	414141	414503	450310		450897	451467
	414117	414464	415714	450681	450820	451208	451288
	433	434	435	481	482	483	484
				_			

Borrelia burgdorferi - Coding regions containing know proteins

	1170	1497	904	289	570	210	209
	66	001	86	96	100	96	66
DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	458681 emblZ12165IB B.burgdorferi gyrA gene encoding DNA gyrase BGYRAG subunit A (partial)	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes,
	451287 gblU045271	452685 gbIU045271	456237 gbIU04527I	emblZ12165ll BGYRAG	464394 gblU03396l	466958 gblU03396	468033 gblU03396l
		452685	456237	458681	464394	466958	468033
	452456	454181	454315	456228	463825	466650	467437
	485	486	487	488	496	497	498
			-		yanning .	-	1

Borrelia burgdorferi - Coding regions containing know proteins

					complete sequence		
-	499	468167	Į.	468433 gblU03396l	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrIA and rrIB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	86	267
	500	468391	468999	468999 gblU03396l	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	95	386
	501	470714	i	470445 gblM88330l	Borrelia burgdorferi 23S ribosomal RNA gene	100	270
_	502	475597	l	480090 gblU03396l	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	97	131
	535	505532	509017	509017 gblL48488l	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	86	2490
	536	509015	513166	513166 gblL484881	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	6	76
	538	513606	514106		Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	100	82
	539	514120	515229	gblU35450I	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	66	1110
quanting	540	515472	516605	516605 gblU49938l	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase CI inhibitor PKCI (pkci) genes, complete cds	666	1134
-	541	516641	517666	517666 gblL241941	Borrelia burgdorferi immunodominant antigen P39 gene, complete cds	66	1026

Borrelia burgdorferi - Coding regions containing know proteins

8 457	909	1461	1386	453	130	314	1404	009
86	66	66	66	100	86	66	100	100
Borrelia burgdorferi (clone pB46) membrane lipoprotein A (bmpA) gene, 3' end, membrane lipoprotein (bmpB) gene, 5' end	Borrelia burgdorferi immunodominant antigen P39 gene, complete cds	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	Borrelia burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	B.bergdorferi (ZS7) YSCI-like gene	B.bergdorferi (ZS7) YSC1-like gene	B.burgdorferi gene for lipoprotein
518256 gblL35050I	518779 gblL24194l	520316 gblU499381	521734 gblU49938I	522204 gblU49938I	522893 gbiU49938I	534772 embIX78708I BBYSC1	535058 embIX78708l BBYSC1	537144 emblX70826l
	l	520316	521734	522204	522893	534772	535058	537144
517732	518168	518856	520349	521752	522168	535086	536461	536545
542	543	544	545	546	547	559	995	561
			_			-		

Borrelia burgdorferi - Coding regions containing know proteins

	57	786	264	56	805		84		354	1185	912	1104	750	1269	1224	969	712	561
	100	100	100	188	92		100		100	100	66	66	66	<u>00</u>	001	001	86	100
	B.burgdorferi gene for lipoprotein	Borrelia burgdorferi 22 kD antigen	Borrelia burgdorferi 22 kD antigen	Borrelia burgdorferi 22 kD antigen	Borrelia burgdorferi periplasmic substrate-	binding protein homolog (p30) gene, complete cds	Borrelia burgdorferi periplasmic substrate-	cds	Borrelia burgdorferi (clone Bb2.13) phoA fusion protein gene, partial cds	Borrelia burgdorferi fesmid clone 31, complete sequence	B.burgdorferi cell division genes	B.burgdorferi cell division genes	B.burgdorferi ftsW, ftsQ & ftsA genes	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence
BBLA7	537191 emblX70826l BBLA7	gb M90084	537968 gblM900841	538757 gblM90084l	572497 gblU291431		574204 gbIU29143I		gb L31422	gbIU43739I	emblX96685l BBCDG	600153 emblX966851 BBCDG	600932 emblX964331 BBFTSWQA	602173 gblU437391	gblU43739I	gbIU43739I	605041 gblL763031	605599 gblU437391
	537191	537665	537968	538757	572497		574204		586936	597983	599052	600153	600932	602173	603394	604087	605041	605599
	537652	539695	537705	538395	574092		575817		585458	596586	297967	599050	600183	506009	602171	603392	604085	602039
	562	563	564	265	909		209		616	629	630	631	632	633	634	635	636	637
	-	1	1	1					-	 -	T	_	_		-	_		

Borrelia burgdorferi - Coding regions containing know proteins

	100 444	100 480	100 378	100 1770	100 1053	100 957	99 1332	99 453	100 630	99 1221	100 447	100 1350	100 231
606938 emblX966851 B.burgdorferi cell division genes BECDG	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence
38 emblX96685l B BBCDG	607379 gbiU437391 B	607861 gbIU437391 B	608208 gblL763031 Branch	609932 gblL763031 ft.	610982 gbIU437391 B	611917 gbIU437391 B	613246 gblL763031 Ba	613674 gblL763031 Bd ffts ffts	614284 gbiU43739l Ba	615470 gblL763031 Bo	615927 gblL763031 Bd fts ftl	617260 gbiU437391 Ba	617507 gbl 0437391 Bo
	ļ	2	607831 60820	608163 60999	606930 6109	610961 6119				614250 61547	615481 61592		617277 6175(
9 869	9 689	640 60	641 6	642 60	643 60	644 6	645 6	646 61	647 61	648 61	649 61	9 059	651 61
			-	_			V	-					

Borrelia burgdorferi - Coding regions containing know proteins

	68 <i>L</i>	288	1062	348					813	189	249
100	001	001	001	001	001	66	100	100	66	100	100
Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flbB), flhF, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flbB), flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flbE), flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL,
618286 gblU43739l	619068 gblU43739l	619653 gblU43739l	620749 gblU43739l	621136 gblU43739l	621755 gblU43739l	622530 gblL759451	gbIL75945I	gbIL759451	623623 gblL759451	622819 gblL759451	623458 gblL759451
]	619068	619653	620749	621136	621755	622530	621822	622802	623623	622819	623458
617498	618280	619066	619688	620789	621114	621742	622028	622515	622811	623007	623706
652	653	654	929	959	159	658	629	099	199	999	663
								_	_		I

Borrelia burgdorferi - Coding regions containing know proteins

	1134	2109	1173	816	345	489	1935	286	78	2439	274	542	327	327
	66	100	100	66	100	100	100	001	100	66	100	66	001	100
fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhF, flbE genes	Borrelia burgdorferi fesmid clone 31, complete sequence	B.burgdorferei promoter element DNA	Borrelia burgdorferi (strain B31) protease (lon) gene, complete cds	Borrelia burgdorferi (strain B31) protease (lon) gene, complete cds	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	Borrelia burgdorferi P1G histone-like protein HBbu (hbb) gene, complete cds					
	624741 gblL759451	626843 gblL759451	628013 gblU43739l	628912 gblU43739l	628807 gbIU43739I	629398 gbIU43739I	631305 gblU43739l	631634 gblU43739l	635476 gblM28682l	649420 gblL772161	gbIL772161	672412 gbIU35673I	672744 gblU35673I	673083 gbiU486511
		626843	628013	628912	628807	629398	631305	631634	635476	649420	649409	672412	672744	673083
	623608	624735	626841	627998	629151	628910	629371	631314	636891	646982	09/159	671567	672418	672751
	664	999	999	<i>L</i> 99	899	699	029	671	9/9	289	889	711	712	713
		-		1	I	Ī	-	-	-	1	-	1	1	-

Borrelia burgdorferi - Coding regions containing know proteins

673491 gblU356731 675118 chi1356731
6/31
675424 gblU35673l
723770 gblU629011
724181 gblU629011
724164 gblU629011
_
gblU629011 Borrelia burgdorferi thdF gene, partial cds, putative motility protein (flbF), flagellar hook associated proteins FlgK (flgK) and FlgL (flgL) genes, complete cds
729308 emblX95669 B.burgdorferi thdF and gidA genes
731176 emblZ12160lB B.burgdorferi thdF, gidA and gidB genes BGIDAG
731799 emblX95668 B.burgdorferi gidA, gidB and moxR genes BBGIDMOX
732848 emblX96434 B.burgdorferi gidB moxR genes and ORF



Borrelia burgdorferi - Coding regions containing know proteins

Borrelia burgdorferi - Coding regions containing know proteins

					genes, complete cds		
	806	857228		858262 gblU28760	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI) genes, complete cds	66	1035
	606	858270	859463	gbIU28760I	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI) genes, complete cds	66	1194
	910	859315	ļ	860226 gblU28760	Borrelia burgdorferi glyceraldehyde-3-phosphate dehydrogenase (GAPDH), phosphoglycerate kinase (PGK), triosephosphate isomerase (TPI) genes, complete cds	66	912
	911	860224	1	860604 gblU576831	Borrelia burgdorferi sequence 3' to the triosephosphate isomerase (TPI) gene	76	183
	912	860645		860316 gblU576831	Borrelia burgdorferi sequence 3' to the triosephosphate isomerase (TPI) gene	95	94
	913	861447	860704	860704 gbIU57684I	Borrelia burgdorferi uracil DNA glycosylase (UDG) gene, partial cds	92	294
	914	861020	861397	861397 gbIU57684l	Borrelia burgdorferi uracil DNA glycosylase (UDG) gene, partial cds	93	244
_	915	861439	862113	862113 gblU57684l	Borrelia burgdorferi uracil DNA glycosylase (UDG) gene, partial cds	96	128
-	930	874089	874859	_	Borrelia burgdorferi 1-acyl-sn-glycerol-3- phosphate acetyltransferase (plsC) gene, 3' end; topoisomerase IV beta-subunit (parE) gene, 5' end	66	408
	931	874877	876679	876679 gblL32861	Borrelia burgdorferi 1-acyl-sn-glycerol-3- phosphate acetyltransferase (plsC) gene, 3' end; topoisomerase IV beta-subunit (parE) gene, 5' end	001	252
	943	887900	886758	886758 emblY088851 BBRUVABH L	B.burgdorferi ruvA, ruvB and queA genes	86	293

Borrelia burgdorferi - Coding regions containing know proteins

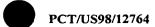
<u> </u>	ł I
890271 emblY088851 B.burgdorferi ruvA, ruvB and queA genes BBRUVABH	9615 890271 embly BBRI L
	719 892404 embl BBPI
	892893 893909 embl)
895371 emblX974491 B.burgdorferi priA and udk genes BBPRIAUDK	973 895371 embly BBPF
895991 emblX974491 BBPRIAUDK	308 895991 embl ³ BBPR
895988 emblX97449l BBPRIAUDK	7976 895988 embl
897963 emblX974491 B.burgdorferi priA and udk genes BBPRIAUDK	i
898555 emblY091411 B.burgdorferi truA gene BBTRUA	899298 898555 embly BBTF

TABLE 3.Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

Contig ID	ORF ID	Start (nt)	Stop (nt)
1	1	2330	1134
1	2	3317	2934
1	8	11375	13021
1	9	11673	11386
1	10	12925	13629
1	11	13538	14146
1	17	25212	24700
1	18	25782	25357
1	19	26115	25870
1	21	27308	27051
1	22	29628	30458
1	29	40696	41217
1	30	41201	41992
1	31	42542	41985
1	32	42593	42982
1	34	44234	44031
. 1	38	48041	47079
1	41	49318	49617
1	43	53234	51810
1	50	59737	58208
1	58	68227	67733
1	65	79757	80404
1	66	81516	80401
1	75	89552	88353
1	82	93338	92766
1	85	95207	95854
1	104	108788	108621
1	105	109764	108943
1	108	112003	111599
1	113	114317	115846
1	114	114522	114316
1	119	118439	118927
1	121	119802	119599
1	125	125688	123967
1	129	128594	129235
1	135	136116	135259
1	136	136558	136298
1	139	139149	139559
. 1	141	140573	140121
1	143	141738	141412
1	145	142218	142060
1	146	142686	142342
1	154	150528	149074
1	158	153832	153981
1	163	158277	158474

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1	171	168052	166205
1	176	171592	171038
1	186	179607	180089
1	189	182345	182046
1	191	182567	182773
1	199	192561	192716
1	205	196592	197476
1	218	207717	206752
1	219	207733	208437
1	221	209337	208915
1	222	209712	209335
1	231	217179	216025
1	238	223660	223418
1	240	224720	225724
1	242	227006	227275
1	248	231761	231501
1	251	232973	233308
1	252	233669	234004
1	254	235115	235456
1	258	241824	242198
1	261	248009	247773
1	269	256846	255872
1	276	265430	265158
1	279	266582	266298
1	281	268474	268280
1	286	274157	274384
1	292	280495	280274
1	294	281344	281042
1	298	287276	285714
1	303	292943	292644
1	304	293273	293037
1	305	294965	294648
1	308	299427	298699
1	309	299051	299212
1	326	320375	319785
1	327	320425	321036
1	331	324198	324413
1	339	332785	332459
1	341	333503	334138
1	342	334116	334739
1	343	334880	335446
11	350	342916	342443
1	351	344789	342897
1	363	357596	356931
1	367	361065	360859
1	370	362519	362196
			502170



Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1	374	366905	366114
1	377	368632	369537
1	378	369928	370560
1	379	370532	371353
1	382	375028	373193
	383	375102	375542
1	387	378677	378198
1	400	394952	394722
	401	396247	394937
1	403	397569	398327
1	406	399103	399294
1	436	416160	416570
1	445	424660	423950
1	446	425181	424642
1	450	428559	428200
1	451	428933	428619
1	455	432590	431628
1	461	437823	438092
1	463	438690	438313
1	466	440749	440222
1	470	441568	441350
1	471	442039	441614
1	472	442216	442037
1	473	442666	442262
1	476	445202	445017
1	493	462106	462519
1	494	462893	462549
1	504	482111	481035
1	505	481552	481800
1	509	483249	483668
1	512	484864	485157
1	516	489171	488527
1	519	492989	492375
1	520	493626	492997
1	521	494169	494864
1	524	497185	497385
1	525	497674	499254
1	527	500251	501294
1	528	501281	502156
1	558	533912	533667
1	568	541267	541491
1	571	544436	544257
1	572	544565	545068
1	578	549603	551198
1	580	551508	551657
1	581	552337	551513

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1			557271
1		561342	561139
1	I	561825	561520
1	592	562536	563360
1	596	565758	566519
1	599	568389	568682
1	602	568680	568856
1	605	570829	571167
1	609	576170	577093
1	612	581549	581091
1	614	582910	584013
1	619	589384	588674
1	624	592665	593465
1	626	594542	595405
1	672	631642	632175
1	677	636650	636892
1	678	637059	638078
1	681	640861	640412
1	686	644887	645207
1	689	649716	649961
1	690	650436	650735
1	691	650733	651056
1	693	653303	653689
1	705	664733	664918
1	707	665979	666770
1	718	679155	678391
1	721	680664	681047
1	722	681523	681849
1	724	681809	682171
1	727	682853	683272
1	734	687648	688067
1	739	691613	692290
1	751	707290	707718
1	763	719197	718904
1	764	720030	719257
1	769	722198	722482
1	783	733736	734647
1	785	735554	736618
1	787	737124	739184
1	792	742924	744801
1	799	753128	752655
1	811	766129	765980
1	812	766438	767772
1	815	770062	769790
1	818	771890	772282
1	831	788219	788836

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

1 838 793566 793414 1 840 794295 794119 1 844 796774 796586 1 852 803096 802908 1 858 809371 809970 1 864 816108 816497 1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 879 831328 83005 1 880 831698 833005 1 880 831698 833005 1 880 831698 833005 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148				
1 840 794295 794119 1 844 796774 796586 1 852 803096 802908 1 858 809371 809970 1 864 816108 816497 1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 849594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109		832		789615
1 844 796774 796586 1 852 803096 802908 1 858 809371 809970 1 864 816108 816497 1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 880 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 923 865660 865346 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139				793414
1 852 803096 802908 1 858 809371 809970 1 864 816108 816497 1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 923 865660 865346 1 923 865660 865346				794119
1 858 809371 809970 1 864 816108 816497 1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273				796586
1 864 816108 816497 1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 928 871012 872580 1 933 878576 879166	1		803096	802908
1 865 816672 817283 1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166				809970
1 866 817281 817838 1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 891 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268	1	864	816108	816497
1 872 823841 824836 1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 949 892388 892924	1			817283
1 876 828191 828739 1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296	1		817281	817838
1 877 828749 829147 1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296	1	872	823841	824836
1 879 831328 831714 1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		1	828739
1 880 831698 833005 1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 923 865660 865346 1 925 868212 869273 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510			828749	829147
1 885 836201 835677 1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510				831714
1 890 841171 840590 1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		831698	833005
1 891 840594 840860 1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		836201	835677
1 899 849453 850148 1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		841171	840590
1 902 851608 852687 1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510				840860
1 918 862867 863109 1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510			849453	850148
1 920 864292 864705 1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510			851608	852687
1 923 865660 865346 1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		862867	863109
1 925 868212 869273 1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510			864292	864705
1 928 871012 872580 1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		865660	865346
1 933 878576 879166 1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1			869273
1 939 884338 883268 1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		871012	872580
1 940 884999 884325 1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		878576	879166
1 949 892388 892924 1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1	939		883268
1 957 900141 899296 1 958 900534 900139 1 959 901526 900510	1		884999	884325
1 958 900534 900139 1 959 901526 900510	1	949	892388	892924
1 959 901526 900510	1		900141	899296
>00010	1	958	900534	900139
	1	959	901526	900510
	1	962	902383	903258

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

TABLE 4.

40	(nt)		•	markin being manne	% sim	% ident
			acession			
		4	4 gil 146582	beta-lactamase [Escherichia coli]	100	0 86
	2 692		240 gil344797	galactosidase fusion protein [unidentified]	1001	
	3 1575		2093 gil458219	ORF 4 [Borrelia burgdorferi]	96	
	41836	414	.59 gil47453	ribosomal protein S12 [Streptococcus pneumoniae]	92	
	14234	1295	51 bbs 161785	60 kda antigen [Borrelia coriaceae, C053, ATCC 4338, Peptide, 514 aal [Borrelia coriaceae]	88	
	1080	1652	2 gnllPIDle2012 50	ORF-D gene product [Borrelia burgdorferi]	88	8 74
	337	26	gnllPIDle1589 79	26 gnllPIDle1589 orfA gene product [Borrelia burgdorferi]	98	5 75
131	1421	1128	gnIIPIDIe 1604 37	gnllPIDle1604 orfD gene product [Borrelia burgdorferi]	85	5 46
	381	674	674 gil458220	ORF 5 [Borrelia burgdorferi]	85	37
	98152	97367	97367 gil 159 1672	phosphate transport system ATP-binding protein [Methanococcus jannaschii]	84	
2 107	108403	109485	gil882454	fructose 1,6-bisphosphate aldolase [Escherichia colil	8	19
19 4		4754	4754 pirlA34520IA3 4520	29K calcium-binding protein, brain-specific - guinea pig (fragments)	81	
20 9	6084	5791	gnllPIDle2012 (49	ORF-C gene product [Borrelia burgdorferi]	8	72
2 52	49986	49600	pirlA027711R7 MCML	pirlA027711R7 ribosomal protein L7/L12 - Micrococcus luteus MCML	08	19
14 1	3071	8	gil1522636	M. jannaschii predicted coding region MJECS02 [Methanococcus jannaschii]	80	09
29 2	218	409	gil1752736	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w Saccharomyces cerevisiae	80	37
32 2	719	925	gil433720	CDC25 [Homo sapiens]	80	73
00	7	946	6 gil 1522636	M. jannaschii predicted coding region MJECS02	80	

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

99	69	58	57	57	09	42	09	57	52	20	55		58	56	56	20	55	36
61	79	78	78	78	78	78	78	78	7.1	77	77		76	76	76	75	75	75
8239 gnllPIDle2881 glucose epimerase [Bacillus thuringiensis]	outer membrane porin protein Oms28 precursor [Borrelia burgdorferi]	ribosomal protein L11 [Thermus aquaticus thermophilus]	4 orfD gene product [Borrelia burgdorferi]	gnllPDle2532 ORF YDL065c [Saccharomyces cerevisiae]	gnllPIDle2012 ORF-B gene product [Borrelia burgdorferi]	CG Site No. 29739 [Escherichia coli]	gnllPIDle2012 ORF-C gene product [Borrelia burgdorferi]	ORF YDL065c [Saccharomyces cerevisiae]	transfer RNA-Tyr synthetase [Bacillus subtilis]	cellobiose phosphotransferase enzyme II" [Bacillus stearothermophilus]	similar to dihydropryridine-sensitive I-type, skeletal	muscle calcium channel alpha-1 subunit (SP:CIC1_RABIT, P07293) [Caenorhabditis elegans]	unknown [Bacillus subtilis]	(pos:59955997,aa:Met) [Bacillus subtilis]	orfC gene product [Borrelia burgdorferi]	674 pirlC30010IC3 hypothetical ORF-6 protein - Sauroleishmania 0010 tarentolae mitochondrion (SGC6)	H. influenzae predicted coding region HI0491 [Haemophilus influenzae]	nusG [Escherichia coli]
9 gnilPIDle288 24	4735 gil1543076	1218 gil587583	2 gn11PIDIe1604 c	gnllPIDle253.	gnllPIDle2017 48	4943 gil882579		gnllPIDle2532 11	gil143795	080 gil466474	536 gil1017809		2183 gil467376	2 gil1065989	3 gnllPIDIe1589 80	pirlC30010IC: 0010	gil1573470	701 gil396321
10823	473;	121	38742	27177	2966	4943	171	742	23697	24080	536		82183	2	3	6674	32163	51701
107148	4878	51661	39290	27416	2382	5107		503	24917	22722	889		81071	208	909	8488	31639	52261
106	4	55	45	46	4	5	_	2	30	34	1		16	1		6	37	26
7	8	7	4 -	ν.	7	19	78	105	7	9	<u>∞</u>	-	3	11	89	7	7	2

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

62	09	37	55	42	58	57	52	49	56	59	58	53	52	47	54	56	63	53	40
75	75	75	75	75	75	74	74	74	74	74	74	73	73	73	73	73	73	. 73	73
$\overline{}$			ORF 2 [Borrelia burgdorferi]	unknown [Borrelia burgdorferi]	4 orfA gene product [Borrelia burgdorferi]	S-adenosylmethionine synthetase [Staphylococcus aureus]	aspartyl-tRNA synthetase [Thermus aquaticus thermophilus]	hypothetical protein [Synechocystis sp.]	974 gnllPIDle1589 orfA gene product [Borrelia burgdorferi]	9 orfC gene product [Borrelia burgdorferi]	CdsK [Borrelia burgdorferi]	[glycoprotein 120 [Simian immunodeficiency virus]	hemolysin [Serpulina hyodysenteriae]	type-I signal peptidase SpsB [Staphylococcus aureus]	4 unknown [Mycobacterium tuberculosis]	Similar to Seryl-tRNA synthetase [Saccharomyces cerevisiae]	5 ORF YGR248w [Saccharomyces cerevisiae]	NADH dehydrogenase, subunit 5 [Acanthamoeba castellanii]	emml gene product [Streptococcus pyogenes]
414 gil520778	652 gnllPIDIe2012 49	62 gnlIPIDle2012 49	578 gil458217	388 gil520783		gil1020317	gil396501	gil1651962	gnllPIDle1589 79	gnilPIDle1589 84	gil1655798	gil406135	gil511145	.262 gil1595810	gnllPIDle2684 56	gil500705	8 gnilPIDle2436 (81	512 gil562035	079 gil694092
414	1652	62	578	388	684	31693	109871	91103	2974	1253	719	7022	21395	44262	62341	91113	93513	3512	8079
653	2437	856	1153	744		30506	111301	92143	4080	468	396	6810	23695	44789	64881	00868	92803	3697	8519
1	3	1	3	1	1	36	109	101	5	7		10	29	26	73	100	106	4	6
20	20	58	89	117	130	2	7	3	20	36	42	2	7	3	3	3	3	4	7

	7	10001	C1112 OC//1	00401	reverse gyrase Methanococcus iannaschii]	73	
14	3	4280	4438 gil520778	87.10	protein p23 [Borrelia burgdorferi]	73	5,5
19	6	7074	6742 gil1773311	73311	NADH dehydrogenase [Ceanothus cuneatus]	73	36
25	3	2369	2587 gil16	gil1655790	CdsC [Borrelia burgdorferi]	73	3
78	2	176	619 gnllP 50	1Dle2012	19 gnllPIDle2012 ORF-D gene product [Borrelia burgdorferi]	73	50
108	1	2	382 gill5	gil1573074	adhesin B precursor (fimA) [Haemophilus influenzae]	73	41
120	1	26	342 gil1978	78	heat shock protein 70 [Sus scrofa]	73	46
3	64	51644	54013 gil1574437	7	sporulation protein (spoIIIE) [Haemophilus influenzae]	72	51
5	9	2899			myosin heavy chain [Gallus gallus]	72	41
9	31	22140	21799 gil895748		putative cellobiose phosphotransferase enzyme II' [Bacillus subtilis]	72	46
8	8	8812	00	66	Orf1 [Borrelia hermsii]	72	55
10	12	8579	8376 gil536681		ORF YBR257w [Saccharomyces cerevisiae]	72	36
45	2	1440	9	7	ErpB2 [Borrelia burgdorferi]	72	42
7	2	1342	(C)	_	pyruvate kinase [Bacillus stearothermophilus]	71	52
7	31	26272	24911 pirIS5 8522	2185	glycyl-tRNA synthetase - Thermus thermophilus	71	54
7	64	95109	58684 gil459009		similar to multifunctional aminoacyl-tRNA	711	48
					synthetase, especially to the prolyl-tRNA synthetase region [Caenorhabditis elegans]		?
m	99	55240	54275 gil21712	1	ORF1 [Synechococcus elongatus]	71	52
m	104	92345	75		secretion protein SecY (AA 1-482) [Mycoplasma capricolum]	71	42
5	43	25567	4		sodium-hydrogen exchange protein-beta [Oncorhynchus mykiss]	71	50
7	3	1179	4		ORF 1 [Borrelia burgdorferi]	71	09
20	4	2964	2392 gil458217		ORF 2 [Borrelia burgdorferi]	71	47
51	7	984	2066 gil1373144		ErpD [Borrelia burgdorferi]	71	41
4 <u>7</u>	1	251	883 gil145280		ORFI [Escherichia coli]	71	40

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

48	41		48	09	35		21	28	4 48	40	47	47	50	CP	74	46	47	38	26	46	42	37
70	70		70	70	70	Î	0 6	0/	0/ 0/	0/0/	202	70	70	69	69	69	69	69	69	69	89	68
538 splQ06797IRL 50S RIBOSOMAL PROTEIN L1 (BL1).	(AE000012) Mycoplasma pneumoniae, phosphocarrier protein HPr; similar to GenBank	Accession Number A49683, from M. capricolum [Mycoplasma pneumoniae]	CheW protein [Salmonella typhimurium]	glycerol kinase [Sulfolobus solfataricus]	cdc4 gene product which is essential for initiation of	DNA replication in yeast [Saccharomyces cerevisiae]	dei AF gene product [Racillus subtilis]	ORF 5 IBorrelia hurodorferil	Orf2 (Borrelia hermsii)	F01G12.6 gene product [Caenorhabditis elegans]	Var1p [Saccharomyces douglasii]	NADH dehydrogenase (ubiquinone) (EC 1.6.5.3) chain 4 - wheat mitochondrion	orfD gene product [Borrelia burgdorferi]	fructose enzyme II [Rhodobacter capsulatus]	YqgI [Bacillus subtilis]	orfI; product unknown [Borrelia burgdorferi]	P30 [Borrelia burgdorferi]	protein 69 [Mycoplasma hyorhinis]	ND6 (AA 1 - 296) [Podospora anserina]	orfA gene product [Borrelia burgdorferi]	ORF' [Escherichia coli]	adenylate kinase [Paracoccus denitrificans]
50538 spiQ06797IRL 1_BACSU	113744 gil1673757		220	73225 gnllPIDle2839 19	93273 gil836815	123 91167913	35807 gil48808	47976 gil1421734	15904 gil1655860	3173 gil 1255880	37 gil 1236921	3970 pirlS16447IS1 6447	1653 gnlIPIDIe1604 37	360 gil151932		94 gil1663561	4	58 gil150176	87 gil13233	2402 gnllPIDle1589 c	30518 gil473817	30 gil1498049
51233	114025		1684	74775	93500	926	35616	48320	16458	2940	5470	4173	1270	65752	99712	25614	14584	7025	8414	1332	29769	72330
54	116		4	84	107	†	47	65	23	4	∞	C	m	69	114	36	71	12	41	7	35	79
2	7		70 (m	3	4	4	4	9	17	20	57	36	2	3	4	٥	77	77	94	2	7

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

42	51	52	52				54	57	43	48	46	44	56	49	38	42	51	49	41	51	48	37		47
89	89	89	89				89	89	89	89	89	89	19	19	19	19	<i>L</i> 9	29	19	19	129	29		129
hypothetical [Haemophilus influenzae]	hypothetical protein [Bacillus subtilis]	D9461.18p; CAI: 0.15 [Saccharomyces cerevisiae]	coded for by C. elegans cDNA CEESS55F; coded for by C. elegans cDNA vk84a1.3; coded for by C.	elegans cDNA yk78g7.3; coded for by C. elegans	yk78g7.5; coded for by C. elegans cDNA yk78g7.5; coded for by C. elegans cDNA yk84a1.5;	strong s	~ · I	ORF 1 [Borrelia burgdorferi]	Orf1 [Borrelia hermsii]	Orf1 [Borrelia hermsii]	ORF 2 [Borrelia burgdorferi]	L8479.4 gene product [Saccharomyces cerevisiae]	50S ribosomal protein L33 [Synechocystis sp.]	ribosomal protein S21 [Myxococcus xanthus]	TagE [Vibrio cholerae]	unknown [Bacillus subtilis]	1502 gnllPIDle2676 alanyl-tRNA synthatase [Thermus aquaticus thermophilus]	60 kda antigen [Borrelia coriaceae, C053, ATCC 4338, Peptide, 514 aal [Borrelia coriaceae]	orfD gene product [Borrelia burgdorferi]	_	SERA protein [Plasmodium falciparum]	gene required for phosphoylation of olioosaccharides/ has high homology with VIDO61	_	_
6385 gil1574032	gnllPIDle2551 17	86074 gil927711	97364 gil1707057			000	0046 gil458217	06/8 gil458216	gil1655859	3694 gil 1655859	gil458217	1133 gil577175	gil1001264	gil710340	gil460955	gil467420	gnllPIDle2676 07	bbs 161785	gnllPIDle1604	276 gil 1655859	889 gnllPIDle8903	906 gil1752736		gnllPIDle1589
106385	68287	86074	97364				40046	406/8	16520	3694	3254	1133	52558	54051	70114	71150	96502	31941	2967	6276	6889	5906		1817
104748	68895	88992	96519			07.707	40048	41916	1/296	2894	3832	927	52752	54290	89069	70653	94703	30304	3590	5524	6611	4995		1221
104	<u>»</u>	86				22	00 [),C	77	2	9	7	57	65	79	81	0110	42	9	6	01	9		2
7 0	<i>S</i>	m	m			+	1 -	1	0 (_ [29	72	7	20 6	m	m (70	4	12	12	12	17		34

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

-				183			
αγ	6	1317	707	mil/59217	ODE 7 (Bornelia humadanfari)		
2	7 8	1740		770 g114.3021.7	ORF 2 [BOITELIA BURGGOTTETI]	/.9	52
ন	33	28572	27751	gil340613	A 'c' was inserted after nt 369 (=nt 10459 in	99	40
					genomic sequence (M10126)) to correct -1 frameshift probably due to gel compression		
2	73	69021	80669	gil153903	methyltransferase (cheR; EC 2.1.1.24) [Salmonella typhimurium]	99	42
7	93	93739	94524	.524 gil45713	P.putida genes rpmH, rnpA, 9k, 60k, 50k, gidA, gidB. uncl and nncB [Pseudomonas mitida]	99	41
3	6	6009	6902	gnllPIDle2639	6902 gnllPIDle2639 OrfD [Streptococcus pneumoniae]	99	47
4	28	20922	20665	1665 gil471731	vacuolating cytotoxin homolog [Helicobacter pylori]	99	50
4	64	47985	47107	107 gil1421735	ORF 6 [Borrelia burgdorferi]	99	43
9	13	7227	8591	8591 gil1591045	hypothetical protein (SP:P31466) [Methanococcus jannaschii]	99	48
34	4	2556	3161	161 gil458218	ORF 3 [Borrelia burgdorferi]	99	42
37	1	985	689	689 gil974334	non-receptor tyrosine kinase [Dictyostelium discoideum]	99	55
3	17	68191	395	gil1651216	Pz-peptidase [Bacillus licheniformis]	65	47
3	123	105911	104070	070 gil 1575784	DNA mismatch repair protein [Aquifex pyrophilus]	65	45
9	6	5726	7126	126 gil1591045	hypothetical protein (SP:P31466) [Methanococcus jannaschii]	65	49
8	6	9684	10325	gnllPIDle2012 50	325 gnllPIDle2012 ORF-D gene product [Borrelia burgdorferi]	65	48
10	I	3	971	gil1373144	ErpD [Borrelia burgdorferi]	65	47
13	5	3956	3411	gil1209872	REV [Borrelia burgdorferi]	65	47
7	92	70509	71069	×	protein-glutamate methylesterase (EC 3.1.1.61) - Salmonella typhimurium	64	45
c	61	48610	50838	gil1001335	soluble lytic transglycosylase [Synechocystis sp.]	64	42
4	5	3519	3773		M protein [Streptococcus pyogenes]	49	32
4	53	38288	37824	824 gil1373141	ORF-10 [Borrelia burgdorferi]	49	50

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30	35	46	30	44	35	41	52	27	49	48	34	43	47	37	45	40	40	38	48	42	45	28
64	64	64	64	64	64	64	64	63	63	63	63	63	63	63	63	63	63	63	63	63	62	69
delta-endotoxin CrylG protoxin [Bacillus thuringiensis]	rhoptry protein [Plasmodium yoelii]	2.9-3 ORF-D [Borrelia burgdorferi]	hypothetical protein [Synechocystis sp.]	gnllPIDle2763 AARP1 protein [Plasmodium falciparum]	P35 antigen protein [Borrelia burgdorferi]	gene required for phosphoylation of oligosaccharides/ has high homology with YJR061w	kinetoplast-associated protein [Trypanosoma cruzi]	2592 gallPIDle2362 ZK287.2 [Caenorhabditis elegans]	carboxyl-terminal protease [Synechocystis sp.]	GLUTAMYL-TRNA SYNTHETASE (EC 6.1.1.17) (GLUTAMATETRNA LIGASE) (GLURS).	TRAB [Plasmid pPD1]	Bts1p [Saccharomyces cerevisiae]	EC 1.1.99.5 [Mus musculus]	glycerol 3 phosphate dehydrogenase [Saccharomyces cerevisiae]	glycerol uptake facilitator [Bacillus subtilis]	ORF-D gene product [Borrelia burgdorferi]	replicative DNA helicase [Bacillus subtilis]	bifunctional protein [Methanococcus jannaschii]	adenine deaminase [Bacillus subtilis]	unknown [Borrelia burgdorferi]	phosphomannose isomerase [Escherichia coli]	cheB peptide [Escherichia coli]
0824 g11402 / 1	gil1041785	gil1209840	gil1652934	gnllPIDIe2763 80	gil1553115	788 gil1752736	gil162142	gnllPIDle2362	gil1652577	266 spIP15189ISY E_RHIME	308 gil 104 l 116	58 gil1098641	237 gil 1339938	gil763191	gil142997	#gnllPIDle2012 (50)	.956 gil467330	853 gil1592217	906 gil633167	268 gil520783	745 gil 146722	573 gil 145524
5 780	4499	19289	2339	839	1177	1788	2	2592	11320	26266	72308	58	71237	71349	74773	4304	24956	3853	9061	268		70573
2985	7798	19738	1608	537	308	1928	589	2837	12750	27753	71067	1056	71398	72845	75552	3747	24123	4161	9558	753	93869	69920
01	7	30	3	-	1	3	1	3	15	32	77	7	82	83	85	9	38	5	13	_	89	75
0	7	7	11	16	19	42	142	7	2	7	7	8	3	ر	3	7	7	11	12	32	7	7

Вотеlia burgdorferi - Putative coding regions of novel proteins similar to know proteins

36	40	37	43	36	35	27	7	36	43	36	32	4	30	45	34	4	48	505	4	38	38	
62	62	62	62	62	69	200	7	62	69	62	- 61	19	61	61	61	61	61	61	61	61	61	
spoOJ93 gene product [Bacillus subtilis]	single-stranded-DNA-specific exonuclease (recJ) [Haemophilus influenzae]	unknown [Helicobacter pylori]	glcB gene product [Staphylococcus carnosus]	glutamine transport ATP-binding protein Q	CigB (Dictyostelium discoideum)	Fu=putative serine/threonine kinase [Drosonhila	melanogaster, Peptide Partial Mutant, 152 aaj [Drosophila melanogaster]	ORF-A gene product [Borrelia burgdorferi]	repeat organellar protein [Plasmodium chabaudi]	ORF-A gene product [Borrelia burgdorferi]	ubiquitin-specific processing protease Saccharomyces cerevisiae1	dnaK homologue [Borrelia burgdorferi]	lipoprotein NlpD [Synechocystis sp.]	YqgP [Bacillus subtilis]	ORF 7 [Borrelia burgdorferi]	CdsJ [Borrelia burgdorferi]	ORF 2 [Borrelia burgdorferi]	ORF 2 [Borrelia burgdorferi]	ORF-D gene product [Borrelia burgdorferi]	methyltransferase [Bacillus aneurinolyticus]	Similar to S. cerevisiae hypothetical protein Ykl012p (Swiss Prot. accession number P33203) and C.	scession number 024600) [Control (Swiss Prot.
5492 gil40031	5212 gil1574144	gil1477770	104 gil 1072419	5144 gil1591493	6976 gil1513302	4378 bbs1144872		8 gnllPIDle1539 (356 gil1151158	gallPIDle1539 (1032 gil 173128	236 gil 143999	gil1653709	261 gil 303863	46478 gil 142 1736	gil1655797	8872 gil458217	gil458217	8652 gnllPIDle2012 50	gil836624	2240 gil1066497	
95492	55212	65677	104	5144	9269	4378		538	356	629	114032	44236	46083	109261	46478	22971	8872	5551	8652	4377	2240	
96334	57341	66414	1762	4431	6743	4563		26	586	138	114352	42737	44821	110052	47119	21496	8300	2006	9398	9079	2449	
3	/9	9/		4	8	9			2		117	55	57	125	63	35		∞	10	12	4	
7 6	2	m	9	<u>∞</u>	19	20		81	106	114	2	3	m	m	4		× į	2	<u>4</u>	15	<u>o</u>	_

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

~				020 BILT0210	Inam-antiporter protein [Enterococcus hirae]	195	32
,	80	70112	6990/	1669 gil 1372995	OrfH (Borrelia burgdorferi)	95	4C
C	116	92686	99212	pirIE22845IE2 2845		56	36
9	26	18732	17791	gil1655797	CdsJ [Borrelia burgdorferi]	95	17
	21	14706	13510	510 gil1574247	H. influenzae predicted coding region HI1410	56	32
	∞	6722	7087	gnllPIDle2428 97	gnllPIDle2428 aBIM [Lactococcus lactis]	56	28
53	7	2446	2018	018 gil1421737	ORF 8 [Borrelia burgdorferi]	75	20
61	2	712	1410	gil583161	albumin binding protein [unidentified]	35	35
7	9	3866	3573	573 gil290487	50S ribosomal subunit protein L28 [Escherichia colil	55	37
7	14	11322	585	gil1303811	YqeU [Bacillus subtilis]	55	33
7	34	28640	82	gil558266	orf gene product [Wolinella succinogenes]	55	30
2	71	69999	67415	gil397486	endonuclease G [Bos taurus]	55	33
3	87	75924	76550	gil403984	deoxyguanosine kinase/deoxyadenosine kinase(I)	55	38
4	99	48434	48958	gil1100900	70 kDa heat shock protein [Theileria parva]	45	32
140	1	322	89	gil15611	gene 17, tail fiber protein [Bacterionhage T7]	55	32
4	34	24244	23867	gil1663563	orfIII; product unknown [Borrelia burgdorferi]	C 2	30
<u>v</u>	6	5510	4179	79 gil1513238	ORFveg 132; similar to Caenorhabditis elegans ORF	54	25
					F59B10.1 encoded by EMBL Accession Number Z49132 [Dictyostelium discoideum]		
<u> </u>	45	27187	25895 ₈	gnllPIDle2614 10	nuclear/mitotic apparatus protein [Xenopus laevis]	54	30
7	28	17905	18162g	62 gil36501	C protein [Homo sapiens]	75	11
	9	4415	5215 _g	5215 gil1707287	putative outer membrane protein [Borrelia	54	25
19	2	1674	2501 g	gil392799	GS/D6 ORF [Dictyostelium discoideum]	24	26
59	ν.	3284	2532 g	gnllPIDle1589 80	orfC gene product [Borrelia burgdorferi]	54	33
31	3	3328	4137p	irlS41649IS4	37 pirlS41649IS4 DNA polymerase - Plasmodium falciparum	1/2	٥٢

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

9 6362 7153 gil1553115 P35 antigen protein [Borrelia burgdorferi] 10 6603 7196 gnllPIDle2563 anti-P falciparum antigenic polypeptide [Saimin schures] 11 12 10333 9422 pir1A427711A4 reticulocyte-binding protein 1 - Plasmodium vivax 12 2771 2771 27711A4 reticulocyte-binding protein 1 - Plasmodium vivax 13 287 gil1498320 Ecli wall-associated protease precursor [Bacillus subtilis] 10 105 106383 107126 gil580905 B.subtilis genes rpmH, rnpA, 50kd, gidA and gidB 11 195 gnllPIDle2202 ps. gene product [Plasmodium falciparum] 11 19 gil182579 CG Site No. 29739 [Escherichia coli] 11 19 100766 101014 gil168365 ord': product uknown [Borrelia burgdorferi] 11 10 100766 101014 gil168365 ord': product uknown [Borrelia burgdorferi] 11 10 100766 101014 gil163365 ord': product lorrelia burgdorferi] 11 10 100766 101014 gil1092864 T03G11.2 gene product [Borrelia burgdorferi] 11 10 100766 101014 gil1092809 ord': gene product [Borrelia burgdorferi] 11 10 100766 101014 gil15353115 P35 antigen protein [Borrelia burgdorferi] 11 10 100766 101014 gil15363 ord': gene product [Borrelia burgdorferi] 12 1277 S04 gil1533115 P35 antigen protein [Borrelia burgdorferi] 10 446 1634 gil1533115 P35 antigen protein [Borrelia burgdorferi] 11 10 6881 7180 gil156218 putative [Cænorhabditis elegans] 11 10 6881 7180 gil156218 putative [Cænorhabditis elegans] 11 12 97006 gil1574476 gadra protein [Sus scord] 11 12 97006 gil172294 protein-tyrosine phosphatase [Saccharomyces		26	34	31	38	25	32	38	31	30	36	27	28	96	32	40	34	22	37	200	77	33
10 6562 7153 gil1553115 9 6362 7196 gallPIDIe2563 93 9422 pirlA427711A4 2771 2771 2771 2771 2771 3 287 gil1498320 91 91 91 91 91 91 91 9		51	51	51	51	51	20	50	20	205	20	50	20	05	50	50	50	49	48	48	48	48
10 6562 7153 gil1553115 9 6362 7196 gallPIDle2563 93 9422 pirlA427711A4 2771 2919 6179 gil173241 1 1 1 1 1 1 1 1 1	jannaschii]	P35 antigen protein [Borrelia burgdorferi]	anti-P.falciparum antigenic polypeptide [Saimiri sciureus]	reticulocyte-binding protein 1 - Plasmodium vivax	ZIP1 protein [Saccharomyces cerevisiae]	cell wall-associated protease precursor [Bacillus subtilis]	B.subtilis genes rpmH, rnpA, 50kd, gidA and gidB Bacillus subtilis]	rps5 gene product [Plasmodium falciparum]	CG Site No. 29739 [Escherichia coli]	103G11.2 gene product [Caenorhabditis elegans]	orfV, product unknown [Borrelia burgdorferi]	VII.1 protein [Streptococcus pyogenes]	orfE gene product [Borrelia burgdorferi]	235 antigen protein [Borrelia burgdorferi]	021 [Borrelia afzelii]	ORF-D gene product [Borrelia burgdorferi]	inknown [Saccharomyces cerevisiae]	TARP antigen [Plasmodium falciparum]	utative [Caenorhabditis elegans]	edA protein (dedA) [Haemophilus influenzae]	astric mucin [Sus scrofa]	protein-tyrosine phosphatase [Saccharomyces cerevisiae]
4 9 6362 10 6603 11 12 10333 12 10333 13 1 14 100766 10 15 106383 10 16 100766 10 17 100766 10 11 100766 10 11 100766 10 11 100766 10 11 10076 10 11 339 10 10 6881 7 10 6881 7 11 97006 96 11 97006 96 14743 144 144		gil1553115	gnllPIDle2563 93	pirlA427711A4 2771	gil173241	gil1498320	gil580905		gil882579	gil1086864	gil1663565	gil49402	gnliPIDle1589 81	gil1553115	gnllPIDle2682 43	gnllPIDle2012 50	gnllPIDle2369 01	gil499325				
3 3 5 7 7 8 8 7 7 8 9 10 10 10 10 10 10 10 10 10 10 10 10 10		7153	7196	9422	6179	287	107126	195	1653		2992	3470	4612	504	1634	941	4	2630	7180	99059	743	14970
3 3 3 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		6362	6603	10333	5919	m	106383	1	50808	100766	23555	4168	5190	1277	1948	582	339	2001	6881	65683	90026	14743
40 1 61 62 62 62 62 62 62 62 62 62 62 62 62 62		5 (01	12	7		105		62	119	32	œ	<u> </u>	2	3	3	_	3	10	75	112	23
		4	01	11	19	23	7	<u>m</u>	3	CC	4	2	0	-	13	92	148	28	m	m	m	

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

28	30	27	96	37	32	28			32	73	21	37	22	23	23	27	27	31	l
48	48	47	47	47	47	46			46	46	46	46	77	45	45	45	45	44	-
M. genitalium predicted coding region MG422 [Mycoplasma genitalium]	chorismate mutase subunit B [Methanococcus jannaschii]	frameshift [Plasmodium falciparum]	ankyrin 3 [Mus musculus]	type I restriction enzyme [Methanococcus januaschii]	P35 antigen protein [Borrelia burgdorferi]	Four tandem repeats of a DNA-binding domain	terminus of CarD. This protein has been purified and	found to bind in vitro to a promoter region [Myxococcus xanthus]	apolipoprotein N-acyltransferase (cute) Haemophilus influenzael	ribosomal protein S19 [Methanococcus jannaschii]	glutamic acid-rich protein [Plasmodium falciparum]	C41G6.i [Caenorhabditis elegans]	picaudalD protein [Drosophila melanogaster]	M. jannaschii predicted coding region MJ0263	ntegrin homolog - yeast (Saccharomyces cerevisiae)	inknown [Saccharomyces cerevisiae]	Inknown [Saccharomyces cerevisiae]	34G8.4 [Caenorhabditis elegans]	repeat Organellar protein [Dlocmodium ob L. 133]
gil1046137	gil1591322	gnIIPIDIe2202 45	gil710551	gil1592264	gil1553115	gil1022328									pirlS30782lS3 i	gnllPIDle2369	gnilPIDle2369	gnllPIDle2364 F	19 pil1151158
929.	282	1.199	80	95240	9941	9471			77324	25719	8816	3648	15	105909	15465	4852	4	81044	5019
7980	2628	5526	55075	94515	9057	9866			78904	24361	9895	3412	632	09271	14212	3950	258	79020	4075
	4	<u></u>	09	94	11	12			68	36	13	4	-	124 1	17	4	-	06	-
1.1	78	77 -	2	7	4	7			8	9	01	<u> </u>	138	3	4	23	92	6	12
	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 [Mycoplasma genitalium]	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 [Mycoplasma genitalium] 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus 48]	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus jannaschii] 48 8 5526 6677 gnllPIDle2202 frameshift [Plasmodium falciparum] 47	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus jannaschii] 48 8 5526 6677 gnllPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus jannaschii] 48 8 5526 6677 gnllPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47 94 94515 95240 gil1592264 type I restriction enzyme [Methanococcus jannaschii] 47	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus jannaschii] 48 8 5526 6677 gallPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47 94 94515 95240 gil1592264 type I restriction enzyme [Methanococcus jannaschiii] 47 11 9057 9941 gil1553115 P35 antigen protein [Borrelia burgdorferi] 47	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus] 48 8 5526 6677 gallPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47 94 94515 95240 gil1592264 type I restriction enzyme [Methanococcus jannaschii] 47 11 9057 9941 gil1553115 P35 antigen protein [Borrelia burgdorferi] 47 12 9986 9471 gil1022328 Four tandem repeats of a DNA-binding domain 46	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus 48 8 5526 6677 gnllPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47 94 94515 95240 gil1592264 type I restriction enzyme [Methanococcus jannaschii] 47 11 9057 9941 gil1553115 P35 antigen protein [Borrelia burgdorferi] 47 12 9986 9471 gil1022328 Four tandem repeats of a DNA-binding domain known as the AT-hook are found at the carboxy terminus of CarD. This protein has been purified and 46	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil159132 chorismate mutase subunit B [Methanococcus 48 8 5526 6677 gnllPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47 94 94515 95240 gil1592264 type I restriction enzyme [Methanococcus jannaschii] 47 11 9057 9941 gil1553115 P35 antigen protein [Borrelia burgdorferi] 47 12 9986 9471 gil1022328 Four tandem repeats of a DNA-binding domain known as the AT-hook are found at the carboxy terminus of CarD. This protein has been purified and found to bind in vitro to a promoter region [Myxococcus xanthus]	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 4 2628 2825 gil159132 chorismate mutase subunit B [Methanococcus 48 8 5526 6677 gnllPIDle2202 frameshift [Plasmodium falciparum] 47 60 55075 55803 gil710551 ankyrin 3 [Mus musculus] 47 94 94515 95240 gil1592264 type I restriction enzyme [Methanococcus jannaschii] 47 11 9057 9941 gil1552115 P35 antigen protein [Borrelia burgdorferi] 47 12 9986 9471 gil1022328 Four tandem repeats of a DNA-binding domain 46 12 9986 9471 gil1022328 Four tandem repeats of a promoter region 46 12 9986 9471 gil1022328 Four tandem repeats of a promoter region 46 12 9986 9471 gil1022328 Four tandem repeats of a promoter region 46 12 9986 78904 77324 gil1573271 apolipoprotein N-acyltransferase (cute) 46	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus 48 2628 2825 gil1591322 chorismate mutase subunit B [Methanococcus 48 47	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48	11 7980 9293 giil 1046137 M. genitailum predicted coding region MG422 48	11 7980 9293 gil1046137 M. genitalium predicted coding region MG422 48

Borrelia burgdorferi - Putative coding regions of novel proteins similar to know proteins

					•		
1/		5 1/35		2 pirlA427711A4 2771	2142 pirlA427711A4 reticulocyte-binding protein 1 - Plasmodium vivax	44	26
22		7 4179		2827 pi1563812	Y CAP CIVE Company		
31		7 1687		1 200511	Action laevisj	44	20
				2/01 gii 1436931	Cutinase negative acting protein [Fusarium solani f. sp. pisi]	43	
3		7 4086		5186 gil343962	VARI protein [Candida olahrata]	5	
28	,	1 11(496 gil157804	Jaminin B2 chain [Drosophila malanaged]	47	25
28	,	5 2889		1 nirlS30787183	integrin homolog	42	23
				0782	0782	42	18
34		1 209		1234 gil 1655797	Cdel [Barrelia burradowil		
65	(*,)	3 1035		1415 oil1654220	cass [Doireira buigabile]]	42	27
2	=	9544		anlibinio 1220	Valiable major protein 16 [Borrella hermsii]	42	34
		· · · · · · · · · · · · · · · · · · ·		6 Stute 1052	Grithidia fasciculata]	41	26
3	122	104072	ľ	3017 gil1151158	Teneat organellar protein [Dlocmodium 1.1.1		
18	9			6366 gill 501/101	M issued in the second of the	41	20
				+7+1701119	Methanococcus januaschiil	40	20
9	9	4662		3964 gil600448	vari profein (aa 1-330) ICandida utilia		
₹	10	7637			microfilerial chaoth anglesis CIIDS II :	39	24
					sigmodontiel	37	19

Borrelia burgdorferi - Coding regions containing to know proteins

TABLE 5.

Contig	Orf ID	Contig Orf ID Start (nt) Stop	\vdash	nt) match	match gene name	nercent	HSP nt
				acession)	ident	length
2			17	402 gblM90084I	Borrelia burgdorferi 22 kD antigen	100	786
2	21	16672	16	310 gblM90084l	Borrelia burgdorferi 22 kD antigen	100	95
2			17	099 gblM90084I	Borrelia burgdorferi 22 kD antigen	100	6
		17415	17	876 emblX70826IB BLA7	B.burgdorferi gene for lipoprotein	100	
2	24	18522		923 emblX70826lB BLA7	B.burgdorferi gene for lipoprotein	100	009
2	25	18606	İ	emblX78708IB BYSC1	20009 emblX78708IB B.bergdorferi (ZS7) YSC1-like gene BYSC1	100	1404
2	26	18661	20295	emblX78708lB BYSC1	295 emblX78708IB B.bergdorferi (ZS7) YSC1-like gene BYSC1	66	314
2	38	32899	32]	74 gblU49938I	Borrelia burgdorferi potential virulence gene cluster	86	130
					membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C11		
	0				inhibitor PKCI (pkci) genes, complete cds		
7	39	33315	32863	363 gblU49938I	Borrelia burgdorferi potential virulence gene cluster	100	453
					(bmpA), BmpB protein (bmpB), putative protein 4,		
					Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds		
2	40	34718	33333	33 gblU49938l	Borrelia burgdorferi potential virulence gene cluster	66	1386
					membrane proteins BmpC (bmpC) and BmpA)
					(bmpA), BmpB protein (bmpB), putative protein 4,		
					ing for name of the complete control inhibitor PKCI (pkci) genes, complete cds	-	
2	41	36211	34751	51 gblU49938I	Borrelia burgdorferi potential virulence gene cluster	66	1461
					membrane proteins BmpC (bmpC) and BmpA		
					(vinpa), burps protein (vinps), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1		•
	1				inhibitor PKCI (pkci) genes, complete cds		
7	42	36899	36288	88 gblL241941	Borrelia burgdorferi immunodominant antigen P39	66	909
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	457	1026	1134	1110	82	76	2490	131	270	386	209
	86	66	66	66	001	76	86	97	100	95	66
gene, complete cds	Borrelia burgdorferi (clone pB46) membrane lipoprotein A (bmpA) gene, 3' end, membrane lipoprotein (bmpB) gene, 5' end	Borrelia burgdorferi immunodominant antigen P39 gene, complete cds	Borrella burgdorferi potential virulence gene cluster membrane proteins BmpC (bmpC) and BmpA (bmpA), BmpB protein (bmpB), putative protein 4, Mg ion transporter MgtE (mgtE), protein kinase C1 inhibitor PKCI (pkci) genes, complete cds	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	Borrelia burgdorferi membrane protein D (bmpD) gene, complete cds	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	Borrelia burgdorferi RNA polymerase beta subunit (rpoB) gene, complete cds, RNA polymerase beta' subunit (rpoC) gene, 5' end of cds	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrIA and rrIB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	Borrelia burgdorferi 23S ribosomal RNA gene	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile- tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile- tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence
	811 gblL35050	gblL241941	38462 gblU49938	838 gblU35450	1961 gblU354501	901 gblL484881	050 gblL484881	977 gblU03396l	620 gblM88330l	066 gblU03396l	041 gblU03396
	811	37401	38462	39838	40961	41901	46050	74977	84620	99098	87041
	37335	38426	39595	40947	41461	46052	49535	79470	84351	86923	87637
	43	4	45	46	47	49	51	83	84	82	98
	2	7	2	2	2	2	2	2	2	2	2

Borrelia burgdorferi - Coding regions containing to know proteins

87 88424 88116 gblU03396l Borrelia burgdorferi B31 Ala-tRNA (rala), Ile-tRNA (ileT), Ile-tRNA (ile-tRNA (ileT), Ile-tRNA (ile-tRNA	210	570	289	904	1497	1170	180	312
88424 88116 gblU033961 91249 90680 gblU033961 98846 96393 emblZ12165IB BGYRAG 100759 98837 gblU045271 102618 103787 gblU045271 103786 103607 gblU045271	96	100	96	86	100	66	001	001
7 88424 88116 gblU033961 8 91249 90680 gblU033961 6 98846 96393 emblZ12165IB BGYRAG 7 100759 98837 gblU045271 8 102618 103787 gblU045271 9 103786 103607 gblU045271	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile- tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	B.burgdorferi gyrA gene encoding DNA gyrase subunit A (partial)	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit gyrB) and ribonuclease P protein component rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds	Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and
88424 8 91249 6 98846 6 98846 7 100759 102618 10 103786 10	gbiU03396i	gblU03396l	emblZ12165lB BGYRAG	gblU045271	gbIU045271	gbIU04527I	gbIU045271	gblU045271
	88116	90680	96393	98837	102389		103607	104177
88 88 96 96 101 101	88424	91249	98846	100759	100893	102618	103786	103866
	87	88	96	97	86	66	100	101
2 2 2 2 2 2	7	2	2	2	2	2	2	2

Borrelia burgdorferi - Coding regions containing to know proteins

ribosomal protein L34 (rpmH) genes, complete cds Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (mpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds Borrelia burgdorferi 212 DNA gyrase b subunit (gyrB) and ribonuclease P protein component (mpA) genes, partial cds, DnaA protein (dnaA), DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete cds Borrelia burgdorferi fesmid clone 31, complete sequence Sequence Sequence SeplU437391 Borrelia burgdorferi fesmid clone 31, complete sequence SeplU437391 Borrelia burgdorferi fesmid clone 31, complete sequence S	Sib Sib	e cds 100 17	1 148	100 1185	99 912	99 1104	100 213	99 750	100 1269	100 1224	100 696	sZ, 98 712 C	100 561	_
	104424 gblU04527 104764 gblU04527 8597 gblU43739 8666 emblX96685 B BCDG 10767 emblX96685 B BCDG 10614 emblX96433 B BFTSWQA 11546 emblX96433 B BFTSWQA 12787 gblU43739 14701 gblU43739 15655 gblL76303	Inbosomal protein L34 (rpmH) genes, complete Borrelia burgdorferi 212 DNA gyrase b subuni (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA) DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete	Borrelia burgdorferi 212 DNA gyrase b subuni (gyrB) and ribonuclease P protein component (rnpA) genes, partial cds, DnaA protein (dnaA) DNA polymerase III beta subunit (dnaN), and ribosomal protein L34 (rpmH) genes, complete	Borrelia burgdorferi fesmid clone 31, complete sequence	B.burgdorferi cell division genes	B.burgdorferi cell division genes	B.burgdorferi ftsW, ftsQ & ftsA genes	B.burgdorferi ftsW, ftsQ & ftsA genes	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ft orf230, smf, hslVU, flgBCE, fliEFGHI, flbAB genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	
	88 88 89 107 107 117 117 117 117 117 117 117	24 gbIU04527I	64 gblU045271	97 gblU43739l		57 emblX96685IB BCDG								

Borrelia burgdorferi - Coding regions containing to know proteins

444	480	378	1770	1053	957	1332	453	630	1221	447	1350	231	789
100	100	100	100	100	100	66	66	100	66	100	100	100	100
Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fisA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hsIVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi ftsA gene, 3' end of cds, ftsZ, orf230, smf, hslVU, flgBCE, fliEFGHI, flbABC genes, complete cds	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence
993 gbiU43739I	475 gbIU43739I	322 gblL763031	146 gblL763031	21596 gblU43739 	131 gblU437391	23860 gblL76303	88 gblL763031	gbIU43739I	26084 gblL763031	26541 gblL76303	27874 gbIU43739I	21 gblU43739l	28900 gbIU43739I
	18475	18822	20546	21596	22531	23860	24288	24898	26084	26541	27874	28121	28900
17550	17996	18445	18777	20544	21575	22529	23836	24269	24864	26095	26525	27891	28112
22	23	24	25	26	27	28	29	30	31	32	33	34	35
3	3	3	3	3	3	3	3	3	8	8	3	3	3

Borrelia burgdorferi - Coding regions containing to know proteins

789	588	1062	348	642	789	207	288	813	249	1134
100	100	100	100	100	66	100	100	100	100	66
Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhE, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhE, flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flhE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flbE genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flbF, flbE, genes	Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flbF, flbE genes
29682 gbIU43739I	67	363 gblU43739I	<u> </u>	369 gbIU43739I	4	436 gblL75945	116 gblL759451	237 gblL759451	772 gblL759451	55 gblL759451
						32642 3243	33129 334]		34320 3407	34222 35355
						42	43	44	45	46
		, (<i>E</i>	3	8	33	C.	m .		3

Borrelia burgdorferi - Coding regions containing to know proteins

2109	1173	816	345	489	1935	286	70	2439	274		542	327	327	411	1566	106
100	100	66	100	100	100	100	001	66	100		66	100	100	66	66	100
Borrelia burgdorferi flagellar hook protein (flgE), flbD, flagellar motor apparatus (motAB), fliL, fliM, fliZ, flagellar export apparatus (fliPQR, flhB), flbF, flbE genes	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	Borrelia burgdorferi fesmid clone 31, complete sequence	B.burgdorferei promoter element DNA	Borrelia burgdorferi (strain B31) protease (lon)	Borrelia burgdorferi (strain B31) protease (lon)	gene, complete cds	S20, Hbb, OrfH and Rho genes, complete cds	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	Borrelia burgdorferi PIG histone-like protein HBbu (hbb) gene, complete cds	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes. complete cds	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds	Borrelia burgdorferi OrfR gene, partial cds, and S20, Hbb, OrfH and Rho genes, complete cds
7457 gblL759451	38627 gblU437391	39526 gblU437391	9421 gblU43739I	0012 gblU437391	919 gblU437391	42248 gbiU43739I	46090 gblM286821	60034 gblL772161	023 gblL77216i	026 abil 1256721	् ।	∞	697 gblU486511	5	732 gblU356731)38 gb <u>U35673 </u>
	37455 38						47505 46		62374 6002	82181 83						85778 8603
47	48	49	20	51	52	53	58	89	69	60	7 6	56	<u>2</u>	95	96	97
m l	33	3	33	3	3	3	3	m	3	C	, (2	2	3	m	£

Borrelia burgdorferi - Coding regions containing to know proteins

1935			1147 gblU61142l	Borrelia burgdorferi outer membrane porin protein Oms28 precursor (oms28) gene, complete cds	66	789
10037 11002 gblU59487	7 11002	gblUS	94871	Borrelia burgdorferi P35 antigen protein gene, and 7.5 kDa lipoprotein gene, complete cds	100	996
11365 11153 gblU59859	11153	gpIUS	98591	Borrelia burgdorferi strain B31 6.6 kDa lipoprotein gene, complete cds	100	213
11577 12230 gblU59487		gblUS	94871	Borrelia burgdorferi P35 antigen protein gene, and 7.5 kDa lipoprotein gene, complete cds	100	373
12578 13414 gblM85216	13	gbIM	352161	Borrelia burgdorferi 27kD protein antigen gene (p27), complete cds	78	370
13753	I		24511	Borrelia burgdorferi 49kb linear plasmid small 12kDa lipoprotein gene, complete cds	66	243
<u>(</u> I	<u>(</u> I	gblL3]	[427]	Borrelia burgdorferi (clone BbK2.1) phoA fusion protein gene, partial cds	100	169
36	36	gbIU75	1/989	Borrelia burgdorferi decorin binding protein B (DbpB) gene, complete cds	66	329
36351 36929 gbIU75867	36	gbIU7.	12989	Borrelia burgdorferi decorin binding protein B (DbpB) gene, complete cds	66	564
36838 36692 <u> gbIU75867 </u>	36	gbIU758	1298	Borrelia burgdorferi decorin binding protein B (DbpB) gene, complete cds	100	147
37001 37624 gbIU75866	1 37	gbIU7.	19989	Borrelia burgdorferi decorin binding protein A (DbpA) gene, complete cds	93	533
39318	68	gbIU42	25991	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	69	731
43349 42447 gb L23137	42447	gbIL23	1371	Borrelia burgdorferi (27985CT2) OspA gene, 3' end and OspB gene, complete cds	66	903
44228 43347 emblA04009IA 04009	43	emblA 04009		B.burgdorferi OspA gene and 5'flanking region	100	882
44792 44403 gblL19702	44403	gblL19		Borrelia burgdorferi outer surface protein A (ospA) and outer surface protein B (ospB) genes, complete cds	88	370
45198 44758 gblL19702	4	gblL19	17021	Borrelia burgdorferi outer surface protein A (ospA) and outer surface protein B (ospB) genes, complete	89	375

Borrelia burgdorferi - Coding regions containing to know proteins

				cds		
4	62	46440	382	Borrelia burgdorferi outer surface protein A (ospA) and outer surface protein B (ospB) genes, complete cds	88	622
4	<i>L</i> 9	49363	50622 gbiL34016	Borrelia burgdorferi (clone 8) S1 gene, complete cds	66	1260
4	89	50708	_	Borrelia burgdorferi (clone 8) S2 gene, complete cds	66	837
4	69	52203	51655 gblL314231	Borrelia burgdorferi (clone BbK2.14) phoA fusion protein gene, partial cds	66	292
4	70	53018	52488 gblL411511	Borrelia burgdorferi (clone 8) s3 gene, complete cds	66	297
5	1	535	71 gblU606421	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	91	465
2	2	1526	546 gblU606421	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	68	374
ς	4	2395	2129 gblL314251	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	86	135
5	11	6832	6542 gblS66708l	{target sequence for detection of Lyme disease agent} [Borrelia burgdorferi, B31, 30-kb circular plasmid pIP87, Plasmid, 416 nt]	97	290
2	12	7422	6817 gblU44914l	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	87	595
2	13	8167	7565 gblU449141	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	84	147
5	14	9408	8284 gblU449141	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	72	268
5	15	10122	9427 gbIU30617I	Borrelia burgdorferi Bbk2.11 (bbk2.10), complete cds	93	260
5	16	10533	11324 gbiU44912l	Borrelia burgdorferi plasmid cp32-1, erpA and erpB genes, complete cds	93	790
2	17	11590	11330 gblŪ44913I	Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	95	261

Borrelia burgdorferi - Coding regions containing to know proteins

173	143]	552	511	801	579	1075	927	379	596	390	384	354	210	440
96	95	100	100	66	86	94	98	80	82	16	66	66	97	95
Borrelia burgdorferi plasmid cp18, OspE (ospE)	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	B.burgdorferi plasmid, orfÅ, B, C, D, E, & F genes, clone pOMB14 and pOMB17	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF. A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	Borrelia burgdorferi B31 BlyA (blyA) and BlyB
588 gblU425991	808 gblU425991	636 emblX87201lB BBRGABCD	185 emblX87201lB BBRGABCD	788 emblX87201lB BBRGABCD	519 emblX87201IB BBRGABCD	158 emblX872011B BBRGABCD	526 gblU45425i	564 gbiU454221	16 gblU454211	gbIU454211	gblU45426I	146 gblU967141	97 gbIU45426I	176 gbIU96714I
1158	11808	1363(14185	14788	15519	16158	18526	18564	19116	19775	20121		20797	21076
11761	13256	14187	14727	15588	16097	17276	17558	19040	19712	20164	20504	20799	21006	21903
18	19	50	21	22	23	24	25	26	27	- 58 - 78	29	30	31	32
5	S	Λ	Λ .	2	5	2	2	2	5	2	5	0	2	5

Borrelia burgdorferi - Coding regions containing to know proteins

	151	467	286	242	317	381	495	300	435	447	465	374	135
	94	06	93	95	96	95	06	97	66	97	96	86	87
(blyB) genes, complete cds	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF. A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF. A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds
\prod													804 gblL31425i B
					2.	3 23	23				30	31	33577 328
	33	4£	35	36	37	38	30	51	52	53	54	55	29
	0	0	S	ς.	2	S	C	v	0	0	S.	2	S

Borrelia burgdorferi - Coding regions containing to know proteins

	657	1590	1212	510	693	375	437	193	140	362	309	756	675	447	1155	345
	100	86	100	100	66	86	77	80	50	96	100	66	100	100	100	100
	Borrelia burgdorferi B31 outer surface protein C (ospC) gene, complete cds	Borrelia burgdorferi 26 kb plasmid GMP synthetase (guaA) gene, complete cds	Borrelia burgdorferi 26 kb plasmid IMP dehydrogenase (guaB) gene, partial cds	Borrelia burgdorferi 26 kb plasmid IMP dehydrogenase (guaB) gene, partial cds	Borrelia burgdorferi transposase-like protein (tra)	Borrelia burgdorferi transposase-like protein (tra) gene, partial cds	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes. complete cds	Borrelia burgdorferi 16 kb plasmid DNA fragment	Borrelia burgdorferi transposase-like protein (tra) gene, partial cds	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi linear plasmid Ip16 DNA, complete sequence	Borrelia burgdorferi linear plasmid Ip16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence
11010101	9022 gbi UU 1894i	11425 gblL258831	12664 gblU13372l	11686 gblU13372l	3 gbIU85588I	677 gbIU85588I	25847 gbIU454231	746 gblU454241	14087 gblU84396l	17876 gblU85588I	507	19,	5862 gblU43414l	7255 gblU43414l	7467 gb U43414	8735 gbIU43414
0730	0/06	9836	11435	12195	695	1081	25041	1420	14287	18352	2815	3522	2188	6089	8621	9079
71	2	17	18	l9		2	39	7	12	17	-	77	2	4	<u>ν</u>	•
7	5	0	0	9	7	7	7	∞	∞ (∞	6	2	2	2	6	5

Borrelia burgdorferi - Coding regions containing to know proteins

110	603	738	273	27.7	210	8/7	143		290	531	Ċ	<u>CI/</u>	224	1202	519		4 <u>4 1</u>	576	210	
100	100	100	00	000	7 0	0	97		91	66	70	00	88	82	81	70	0/	84	91	
Borrelia burgdorferi linear plasmid lp16 DNA.	complete sequence Borrelia burgdorferi linear plasmid lp16 DNA,	complete sequence Borrelia burgdorferi linear plasmid lp16 DNA,	complete sequence Borrelia burgdorferi linear plasmid lp16 DNA	complete sequence Borrelia burgdorferi linear plasmid ln16 DNA	complete sequence Borrelia hirsdorferi 7 9-7 locus ORE Crease	partial cds, ORF-D, REP+, REP-, and lipoprotein	Borrelia burgdorferi 16 kb plasmid hypothetical	protein gene, complete cds	Borrelia burgdorferi Ip21 circular plasmid, complete sequence	Borrelia burgdorferi exported neurotoxin-like	Borrelia huradorferi In21 circular plasmid	complete sequence	Borrelia burgdorferi Ip21 circular plasmid,	Borrelia burgdorferi plasmid cp18, OspE (ospE)	B.burgdorferi repeated DNA element, 30.5 kb	circular plasmid copy Borrella burgdorferi In21 circular plasmid	complete sequence	Borrelia burgdorferi Ip21 circular plasmid, complete sequence	Borrelia burgdorferi Ip21 circular plasmid,	complete sequence
214 gblU43414l		107 gblU43414l	027 gblU43414l	241 gblU43414l			2886 gblU12332l		_	983 gblL166251 F	001 gblU036411)	167 gblU036411	41 gblU425991	88 emblX87127IB B	55 gblU036411		68 gblU036411	9544 gbiU036411 B	ည
10224 9	10370 10	11844 11	13299 13	13612 13.	2164 10	······································	2686 28			1525	4098 49		4691 44 	6348 50	6673 77	7786 83		8393 89	9290 95.	
7	<u></u>	6	10	111	2		m	-	-	77	9	Ţ		8	6	10		-	12	
6	6	6	6	6	10		10	13	C	13	13	2	13	13	13	13		13	13	

Borrelia burgdorferi - Coding regions containing to know proteins

366	85 242		77 267	95 296		93 594									
s cds	e cds	pD)	v					1.	÷ 0,						
Borrelia burgdorferi protein p23 gene, complete cds	Borrelia burgdorferi protein p23 gene, complete cds	Borrelia burgdorferi outer surface protein D (ospD) gene, complete cds	Borrelia burgdorferi (clone 8) s3 gene, complete cds	Borrelia burgdorferi plasmid cp32-1 PCR target site, partial sequence	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia hirodorferi nlasmid cn32-2 em and	D genes, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene complete cds Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	erpD genes, complete cds Borrelia burgdorferi strain 297CH putative outer membrane protein (ospF) gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp32-4, erpH gene, complete cds Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds
6217 gb L31616 Bo		2854 gblM97452l Bo	657 gblL411511 Bor	4 gbIU609631 Bo	834 gblU44914l Bo	581 gblU44914l Bo									
6217 ₁₈	6671 g	2854 g	3657 g	4	834 g	1581 g		2257 g	2257 g 2964 g	2257 g 2964 g 5143 g	2257 g 2964 g 5143 g 5183 g	2257 g 2964 g 5143 g 5183 g 5360 g	2257 g 2964 g 5143 g 5183 g 5360 g	2257 g 2964 g 5143 g 5183 g 5360 g 317 g	2257 g 2964 g 5183 g 5360 g 317 g 658 g
2168	6126	3660	3136	849	1427	2168		2946	2946	3794	2946 3794 4334 5362	2946 3794 4334 5362 5581	2946 3794 4334 5362 5581 306	2946 3794 4334 5362 5581 664	2946 3794 4334 5362 5581 564 664
9	7	5	CC .	1	2	3		4	4 8	4 0	4 8 9 6	4 2 9 5 8	4 2 9 2	4 2 0 1 8 - 2	4 0 0 6 8 - 2 6
14	14	16	19	21	21	21	_	21	21	21 21 21	21 21 21 21 21 21 21 21 21 21 21 21 21 2	21 21 21 21 21 21 21 21 21 21 21 21 21 2	21 21 21 21 21 21 21 21 21 21 21 21 21 2	22 22 22 22 22 22 22 22 23 24 24 24 24 24 24 24 24 24 24 24 24 24	22 22 21 21 21 21 21 21 21 21 21 21 21 2

Borrelia burgdorferi - Coding regions containing to know proteins

	750	378	204	603	221	362	220	478	309	219	610	419	786	615
	94	100	100	96	96	94	08	87	86	96	86	97	100	100
sedneuce	108 gblU76406l Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence		536 gblU43414l Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	82 emblX87127IB B.burgdorferi repeated DNA element, 30.5 kb BPBRGEA circular plasmid copy	~	873 gblAF000270l Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	2621 gblU454271 Borrelia burgdorferi 2.9-7 locus, ORF-A-D, REV, and lipoprotein (LPA and LPB) genes, complete cds						45 emblX87127IB B.burgdorferi repeated DNA element, 30.5 kb BPBRGEA circular plasmid copy	71B
	5108 gbi	760 gbl	1536 gbl	82 em BP	682 em BP	2573 gbl.	2621 gbl	149 gbl	4355 gblī	434 gblL316151	258 gblI	686 emt BPF	1545 emb BPE	
		383	1333	684	2				4		7			1543 21
		•				7	3(37	46	ς,	13	7	7	15.
	2		2		2	4	5	9	∞		7	7	7	3
	24	25	52	56	56	97	26	50	97	17	27	ر ا	9	30

Borrelia burgdorferi - Coding regions containing to know proteins

	100 645	92 976	100 546	100 1152	98 379	93 577	91 174	100 240		92 571	100 133	100 534	100 597	99 529	971 170
	30.5 kb	pE (ospE)			rpC and	rpC and	aT), Ile- rlA and nes,			pC and				(Đ)	
circular plasmid copy	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	Borrelia burgdorferi plasmid cp32-5, erpl gene, complete cds	Borrelia burgdorferi plasmid cp32-1, erpA and erpB2 genes, complete cds	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia burgdorferi B31 Ala-tRNA (alaT), Ile-tRNA (ileT), 16S rRNA, 23S rRNA (rrlA and rrlB), and 5S rRNA (rrfA and rrfB) genes, complete sequence	Borrelia burgdorferi 23S ribosomal RNA gene	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia burgdorferi plasmid cp32-3, ErpG (erpG) and BapA (bapA) genes, complete cds	B.burgdorferi ospG and bapA genes	1604 emblX82409IB B.burgdorferi ospG and bapA genes BOSPG	Borrelia burgdorferi plasmid cp32-3, ErpG (erpG) and BapA (bapA) genes, complete cds	B.burgdorferi plasmid, orfA, B, C, D, E, & G
BPBRGEA	7IB		995 gbIU72996I		882 gblU44914l			737 gblM88330l		28 gblU44914l E	653 gblU42598l E	983 emblX82409IB B BOSPG	1604 embiX82409IB B BOSPG		665 emblX87202 B
	2158	3247	450	1008	2253	3050	co.	976	-	672	850	1516	2200	2602	296
	4	5		2	<i>c</i>	4		7			2	3	4	<u>ν</u>	
-	30	30	33	33	33	33	35	35	36	38	38	38	38	38	36

Borrelia burgdorferi - Coding regions containing to know proteins

BBRGBCDE genes, clone pOMB10 253 gblU42599 Borrelia burgdorferi plasmid cp18, OspE (ospE) 91 2572 emblX87201IB Burgdorferi plasmid, orfA, B, C, D, E, & F 94 BBRGABCD genes, clone pOMB14 and pOMB17 2572 emblX87127IB Burgdorferi repeated DNA element, 30.5 kb 91 2572 emblX87127IB Burgdorferi repeated DNA element, 30.5 kb 93 2586 emblX87127IB Burgdorferi repeated DNA element, 30.5 kb 93 2586 emblX87127IB Burgdorferi repeated DNA element, 30.5 kb 93 2586 emblX87127IB Burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence sequenc	06
	OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like
2284 2284 2572 2861 2861 1732 1732 1747 1747 1747 17418	
2353 2574 2874 2874 3000 3000 342 1172 1173 1133 635	
6 4 8 0 - 2 6 - 2 6 - 2 - - 2 -	
39 39 39 39 40 40 40 40 41 41 41 43 43	

Borrelia burgdorferi - Coding regions containing to know proteins

1 1				orf1 gene, partial cds		
<i>x</i>	2242	1784	1784 gblU45423I	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	85	421
4	2860	2318	2318 gblU454211	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	259
	1158	178	178 gbIU60642I	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	89	374
3	2531	1761	1761 gblL314251	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	66	135
	287	3	3 gbIU78764	Borrelia burgdorferi plasmid cp32-1, erpA and erpB2 genes, complete cds	84	153
	2037	1453	1453 gblL13924l	Borrelia burgdorferi outer surface protein E (OspE) gene, complete cds	06	386
₹	2663	2893	2893 gblU44912l	Borrelia burgdorferi plasmid cp32-1, erpA and erpB genes, complete cds	06	230
	174	338	338 gbIU425991	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	96	91
7	259	996		Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	100	692
	964	1527		Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	100	564
4	1509	2111	2111 gblU425991	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	66	603
2	2537	2851	emblX872011B BBRGABCD	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	86	315
	2	526	526 gblU45425i	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	95	525
	1245	724	724 gblU454241	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	94	483

Borrelia burgdorferi - Coding regions containing to know proteins

651	327	91	804	909	1596	612	269	146	140	146	422	489	101
68	87	100	66	66	86	66	98	94	94	98	81	66	001
Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia burgdorferi plasmid cp32-6, erpK gene, complete cds	Borrelia burgdorferi plasmid cp32-6, erpK gene, complete cds	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	B.burgdorferi plasmid, orfA, B, C, D, E, & G genes, clone pOMB10	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like
1321 gblU45424 	25 gbIU44914I	1182 gb U72997I	1244 gbIU72997I	18 gblU764061	704 gblU764061	7	7	236 emblX87202lB BBRGBCDE	6	0	1	581 gblAF0002701	719 gblAF000270l
1971	363	412	2047	713	2308	613	2203	3		250	1650	93	883
3	1	2	<u>e</u>		2		3	1	2	6	9		2
47	48	48	48	49	49	51	51	52	52	52	52	53	53

Borrelia burgdorferi - Coding regions containing to know proteins

			orf1 gene, partial cds		
	811	811 gblAF000270	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	100	289
	1064	064 gblAF000270	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	96	381
	1380	380 gbIU45427I	Borrelia burgdorferi 2.9-7 locus, ORF-A-D, REV, and lipoprotein (LPA and LPB) genes, complete cds	93	362
	1740 ₺	40 gblU45426I	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	86	210
	434 8	134 gbIU45422I	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	92	326
	471	71 gblAF000270	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	86	362
	2109 el B	09 emblX87127IB BPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	84	246
	1800 g	gblL314251	Borrelia burgdorferi (clone BbK3.168) phoA fusion protein gene, partial cds	06	118
	1111e B		B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	75	519
	694 e B	694 emblX87127IB BPBRGEA	B.burgdorferi repeated DNA element, 30.5 kb circular plasmid copy	72	786
	1410e B		B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	95	498
	3e B		B.burgdorferi plasmid, orfÅ, B, C, D, E, & F genes, clone pOMB14 and pOMB17	62	260
-	282 er	82 emblX87202lB	B.burgdorferi plasmid, orfA, B, C, D, E, & G	74	501

Borrelia burgdorferi - Coding regions containing to know proteins

	351	510	204	300	435	440	207	384	390	342	374	393	281	552
	78	100	100	93	96	94	86	66	86	66	86	96	85	91
genes, clone pOMB10	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	Borrelia burgdorferi linear plasmid Ip16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes, complete cds and REP+ gene, partial cds	Borrelia burgdorferi 2.9-3 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-5 locus, ORF-A-D,
BBRGBCDE	910 gblU42599I	gblU43414I	117 gblU43414l	75 gbIU60642I	641 gbIU60642I	018 gblU60642l	gblU96714l	600 gblU454261	946 gbIU454231	083 gblAF000270I	gbIU60642I	gbIU60642I	12 gblU45422l	340 gblU45425I
	910	54	1117	75	641	1018	275	009	946	1083	925	1328	12	540
	1704	563	1320	647	1075	1530	33	217	557	1424	7	936	464	1256
-	m		7		77	m		7	Č.	4		2		ন
-	62	49	49	99	99	99	70	0/	70	70	75	75	9	76

Borrelia burgdorferi - Coding regions containing to know proteins

	379	651	255	1198	347	440	151	486	148	135	195	243	447
	06	76	08	66	91	84	08	98	97	67	86	86	86
cds	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-4 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	Borrelia burgdorferi B31 BlyA (blyA) and BlyB (blyB) genes, complete cds	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-7 locus, ORF-A-D, REV, and lipoprotein (LPA and LPB) genes, complete cds	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at
	2 gblU454221	509 gblU45424l)34 gbIU43414I	202 gblU764061	360 gblU454211	gblU967141	gblU967141	289 gblU454221	54 gblU45427I	gbIU60642I	gbIU60642I	23 gbl 0606421	08 gblU60642I
	7	509	1034	1202	360	1008	636	289	954		31	323	508
	433	1159	657	က		358	791	891	1151	137	325	565	954
-		7	7			7	<i>c</i>		21		7	<u>~</u>	4
100		<u> </u>	81	83	8	82	85	98	98	<u>8</u>	8 8	<u> </u>	<u>×</u>

Borrelia burgdorferi - Coding regions containing to know proteins

201	313	. 331	368	243	458	472	380	234	220	234	477	889	146
86	97	68	96	76	06	94	70	100	86	100	66	66	100
Borrelia burgdorferi plasmid cp32-2, sequence at position 5kb	Borrelia burgdorferi 2.9-2 locus, ORF-C gene, partial cds, ORF-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi strain B31 2.9-like locus, OrfC, OrfD, Rev (rev), lipoprotein (LP), and 36 kDa-like orf2 genes, complete cds, and 36 kDa-like orf1 gene, partial cds	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia burgdorferi plasmid cp18, OspE (ospE) gene, partial cds	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi linear plasmid Ip16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi putative vls recombination cassettes Vls2-Vls16b (vls) gene, complete sequence	Borrelia burgdorferi 2.9-6 locus, ORF-A-D genes,
891 gbIU60640I	34 gbl∪45422l	gblU454211	940 gbIU454211	245 gblU45425I	282 gblAF000270	3 gbIU44914 	264 gblU425991	408 gb U43414	757 gblU43414l	440 gblU43414l	837 gblU43414l	911 gb U76406	242 gblU45426l
891	34	578	940	245	282	3	264	408	757	440 8	837	91118	242 g
1091	927	162	572	co .	749	506	827	175	329	207	361	3	388
5		_	7	 -	79	—	-		7	-	7		1
88	91	93	93	94	94	97	86	66	66	I0I	101	102	104

Borrelia burgdorferi - Coding regions containing to know proteins

210	789	2001	306	228	456	310	405	300	374	391	234	356	238	234
100	95	70	93	66	95	93	68	86	86	85	100	66	100	1001
complete cds and REP+ gene, partial cds Borrelia burgdorferi B31 BlyA (blyA) and BlyB	(blyB) genes, complete cds Borrelia burgdorferi 2.9-5 locus, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete	cds Borrelia burgdorferi protein n/3 gene complete cds	Borrelia burgdorferi protein p23 gene, complete cds	Borrelia burgdorferi (clone BbK2.5-6) unknown protein gene, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi 2.9-1 locus, ORF 5-8, ORF-A-D, REP+, REP-, and lipoprotein (LP) genes, complete cds	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-4, sequence at position 4-6kb	Borrelia burgdorferi plasmid cp32-2, erpC and erpD genes, complete cds	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	Borrelia burgdorferi linear plasmid lp16 DNA, complete sequence	B.burgdorferi plasmid, orfA, B, C, D, E, & F genes, clone pOMB14 and pOMB17	Borrelia burgdorferi linear plasmid lp16 DNA,
386 gblU967141	gbIU45425I	4 gblL316161	173 gblL316161	580 gblL316151	456 gbIU454211	761 gblU454211	215 gblU45421	84 gbIU60642I	23 gblU60642I	gblU44914I	gblU43414I	'00 gbIU43414I	697 emblX87201IB BBRGABCD	467 gblU43414
386	811	4	173	280	456	761	215	84	123	2	408	700	697	467
595	2	264	298	807		450	787	653	719	403	175	329	458	234
72			2	8		7	_	-	= -	=-	-	2		- -
104	107	109	109	109	110	0110		119	171	77.1	87	78	671	132

Borrelia burgdorferi - Coding regions containing to know proteins

171		243	331	513	1	153)	432)	495	}	144		706	<u> </u>	351		
66		08	78	100)	100		86		94	`	98)	88	}	16		
Borrelia burgdorferi linear plasmid lp16 DNA,	complete sequence	309 embl x 8 / 12 / 1B B. burgdorferi repeated DNA element, 30.5 kb BPBRGEA circular plasmid conv	B.burgdorferi plasmid, orfA, B, C, D, E, & G	Borrelia burgdorferi B31 BlyA (blyA) and BlyB	(blyB) genes, complete cds	Borrelia burgdorferi B31 BlyA (blyA) and BlyB	(blyB) genes, complete cds	Borrelia burgdorferi plasmid cp32-3, ErpG (erpG)	and BapA (bapA) genes, complete cds	3 emblX87127IB B.burgdorferi repeated DNA element, 30.5 kb	circular plasmid copy	Borrelia burgdorferi Ip21 circular plasmid,	complete sequence	Borrelia burgdorferi plasmid cp32-4, sequence at	4-6kb	Borrelia burgdorferi GrpE protein homologue gene.	DnaK protein homologue gene, and DnaJ protein	homologue gene, complete cds's
Borreli	comple	B.burge circular		Borrelia	(blyB) (Borrelia	(blyB) g	Borrelia	and Bar	B.burgo	circular	Borrelia	complet	Borrelia	position 4-6kb	Borrelia	DnaK pi	homolog
660 gblU434141	GIEGI FOXI	emblX8/12/IB BPBRGEA	4 emblX87202lB BBRGBCDE	33 gblU967141		276 gblU96714I		498 gbIU42598I		emblX87127IB	BPBRGEA	2 gblU036411		542 gbIU60642I		2 gblM96847I		
099	003	000	4	33		276		498		m		7		542		ন		
388	,	n	339	554		124		19		497		193	1	m		352	-	
7	-	-	-		-	7							-	=	-	_		
132	133	133	134	141		141	- (143	-	144		146		147		133		
																•		_

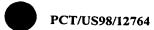


TABLE 6.

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

Contig ID	ORF ID	Start (nt)	Stop (nt)	
2	4	2730	35	54
2	5	3559	34	
2	7	5464	38	
2	13	10502	99	
2	17	13800	135	
2	19	15368	152	
2	28	21155	214	
2	50	41944	421	86
2	58	53786	529	11
2	59	54816	537	73
2	61	57393	558	13
2	63	57882	576	82
2	65	60898	602	
2	66	61441	620	
2	67	62078	626	
2	70	65896	6654	
2	74	70203	699	
2	78	71818	7139	
2 2	80	72956	740:	
	81	73515	7320	
2 2	90	92181	9252	
2	91	92968	9255	
2	108	109872	11005	
2	112 113	112408	1128	
2	113	112858 113035	11303	
2	115	113506	11346	
2	119	114325	11372 11485	
3	6	3279	407	
3	8	5156	601	
3	54	42256	4278	
3	59	47264	4750	
3	60	47673	4869	
3	63	51475	5102	
3	70	60330	6057	
3	71	61050	6134	
3	72	61347	6167	
3	74	63917	6430	
3	86	75347	7553	
3	88	76593	7738	
3	99	89769	8900	
3	102	91278	9166	
3	103	92137	9246	
3	105	92423	9278	
3	108	93467	9388	
3	115	98262	9868	

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

3	121	102227	10200
3	126	111308	102904
4	6	3751	110055
4	7	4218	4179
4	19	16115	5042
4	20	17028	15516
4	21	17379	16075 17092
4	22	17735	17092
4	24	19243	18785
4	25	18942	19196
4	26	20677	19259
4	27	19431	19751
4	29	21376	20876
4	30	21899	21423
4	31	22918	21845
4	33	23951	23553
4	37	26253	25627
4	38	26991	26332
4	39	28181	26931
4	40	29175	28522
4	43	30605	30342
4	45	34906	33548
4	48	35750	35932
5	3	2102	1527
5	5	2656	2393
5	7	3460	2900
5	10	6544	5645
5	40	25278	24322
5	41	25235	25600
5	42	25665	25276
5	44	25881	25663
5	48	27883	27410
5	49	28351	27881
5	50	29028 29454	28324
5	56	32199	29026 31666
5	57	32571	32200
5	58	32826	32569
5	60	32913	33245
5	61	33766	33575
5	62	34173	33742
5	64	35514	34861
6	2	954	1181
6	3	1590	1763
6	5	3400	3954
6	7	4691	5218
6	8	5187	5699

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Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

		·	
6		6498	
6	12	6975	6727
6	14	7978	7448
6	15	8479	7976
6	22	15106	15636
6	27	19999	18842
6	28	20036	20668
6	29	21814	20690
6	30	20949	21269
6	35	24136	23630
6	37	25697	26248
7	8	8100	7792
7	10 11	8145	8288
7		9374	8517
7	12 13	9771	9325
7	13	9652	10185
7	15	10163 10517	9765
7	16	11363	10173
7	17	11363	10524
7	18	12495	11392
7	19	13516	11902
$\frac{1}{7}$	20	12807	12473 13154
7	22	15149	13134
7	24	15855	15046
7	25	15503	15826
7	26	16638	15853
7	27	19344	16636
7	31	19473	19727
7	32	20067	19675
7	33	20762	20049
7	34	21136	20738
7	36	22975	23406
7	40	26667	25870
8	3	2907	4118
8	5	5898	6059
8	6	7399	8313
8	13	15645	15899
8	14	17281	16331
8	15	16905	17111
10	4	3211	3684
10	6	3857	4456
10	8	5982	5599
10	11	8038	7802
10	14	10255	10100
11	7	5688	5828
11	9	7248	7685

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

11	10	7672	8028
11	13	9642	10154
12	1	101	370
12	2	982	680
12	3	1390	1115
12	4	1528	
12	5	1913	1388
12	11		1431
14	2	7308	6616
14		3588	3328
14	9	4657	4815
15		7981	8511
15	1	1	327
15	2	325	1077
	3	1478	657
15	4	2360	1758
15	5	2839	2507
15	9	3922	3743
15	10	4145	3900
15	11	4112	4270
15	13	7677	6127
15	14	7852	7709
15	15	8052	7825
15	16	8222	7857
16	2	1733	1936
16	3	1905	2063
16	6	5212	4220
16	7	8903	8505
17	2	1500	1709
17	5	4097	4660
17	7	6344	6189
18	1	1635	2465
18	2	2509	3306
18	3	3332	4390
18	5	4933	4727
18	7	6353	7084
18	8	7098	7625
20	7	4700	4557
22	4	2175	1228
22	5	2132	2314
22	6	2829	2173
22	8	3254	3601
22	9	4408	4169
22	10	4875	4402
22	11	5343	4873
23	2	2283	1537
23	3	3564	2617
25	6	3677	4147
	1	30,7	71-7/

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

26		4251	3889
28		732	1739
29	3	310	885
31	1	28	195
32	3	935	1603
32		1637	2332
37	2	1379	
42			1059
44	4	2708	2388
	2	1734	1159
44	4	2942	2532
47	4	2336	2115
50	1	908	120
52	4	674	501
56	1	152	1465
56	2	611	459
56	3	1479	2150
58	3	1691	1329
58	5	1867	2046
59	2	2018	
61			1044
	1	1	657
61	3	1389	1907
62	4	1115	1345
63	1	663	325
63	2	769	446
63	3	1759	1013
65	1	472	903
65	2	901	1236
67	1	387	4
67	2	979	401
67	3	1482	961
68	2		
69	3	451	612
		840	574
71	1	363	4
72	1	586	933
73	1	300	4
73	2 3 1	824	279
73	3	1396	1145
79	1	22	1119
82	1	701	303
82	2	1188	775
84		331	134
84	2	983	
87			348
87	1	277	2
	2	1136	267
96	1	434	57
96	2	748	557
97	2	976	659

Borrelia burgdorferi - Putative coding regions of novel proteins not similar to know proteins

103		301	2
103	L	886	
105	_	36	
106		425	3
106		761	600
112	1	416	799
113	1	685	59
118	1	1	489
118	2	487	753
120	2	299	691
124	1	1	630
127	1	702	322
135	1	287	3
135	2	649	407
136	1	1	645
140	2	619	332
145	1	1	480

(1) GENERAL INFORMATION:

- (i) APPLICANT: Human Genome Sciences, Inc. et al.
- (ii) TITLE OF INVENTION: Borrelia burgdorferi Polynucleotides and Sequences
- (iii) NUMBER OF SEQUENCES: 155
- (iv) CORRESPONDENCE ADDRESS:
 - (A) ADDRESSEE: Human Genome Sciences, Inc.
 - (B) STREET: 9410 Key West Avenue
 - (C) CITY: Rockville
 - (D) STATE: Maryland
 - (E) COUNTRY: USA
 - (F) ZIP: 20850

(v) COMPUTER READABLE FORM:

- (A) MEDIUM TYPE: Diskette, 3.50 inch, 1.4Mb storage
- (B) COMPUTER: HP Vectra 486/33
- (C) OPERATING SYSTEM: MSDOS version 6.2
- (D) SOFTWARE: ASCII Text
- (vi) CURRENT APPLICATION DATA:
 - (A) APPLICATION NUMBER:
 - (B) FILING DATE: Herewith
 - (C) CLASSIFICATION:
- (vii) PRIOR APPLICATION DATA:
 - (A) APPLICATION NUMBER:
 - (B) FILING DATE:
- (viii) ATTORNEY/AGENT INFORMATION:



- (A) NAME: Brookes, A. Anders
- (B) REGISTRATION NUMBER: 36,373
- (C) REFERENCE/DOCKET NUMBER: PB370PCT

(vi) TELECOMMUNICATION INFORMATION:

- (A) TELEPHONE: (301) 309-8504
- (B) TELEFAX: (301) 309-8512

(2) INFORMATION FOR SEQ ID NO: 1:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 910715 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: double
- (D) TOPOLOGY: linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1:

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TTGTATATTC	TATAGAAAAA	ACGATTAGAA	TTAAACAAAG	CCATAACTGA	ACCAACGGTA	120
ATTAGTAGAT	AAAGGGATCA	AAATATTTTT	TATTGCAGCA	AGAATACCTT	GGTATATTAG	180
AAAAACCAAA	AGTCATAGTC	AAATCATCTT	TTGATAACAA	TCCCCAAATC	ТАТААТТТАТ	240
ТАТСАААТТА	ATTGCTCCCT	TGAAAAGATT	AGTTTTTAAA	ACTACAAGAC	ТАСТАТСААТ	300
CACTATCAGA	TAGATTAAAA	CAACCTTTAC	AAGAAAAAA	TCTTACTACT	ATTTTATTGT	360
AAATGTATTA	TAAAATAAGT	TCATGCAAAA	ACTTACAATT	TTTCACAACA	AACTACAATA	420
AAATCATGTA	AACAAACAAT	TTCTTTGAAA	ATTAAGCAAA	ТТТАТАААТА	ТАААТТАТАА	480
AGATATATAT	TTTTATATGA	ТСААТААТАА	AAATTAATAG	GATACTTATT	TGGAAAAATT	540
ATTGAAAAAA	CAATAAGCAT	GAATTGCCAC	AATAAGCTAA	TTGTCACTTA	ATAATTCTTG	600
TTTACTAGAC	CACATTAGTA	TAAACTCAAA	TATTGGCTAC	TATAATATAG	GGGCTTTATA	660
CGCCACATGT	TTAATGATAA	CATAAGAAAA	TATTGCAATA	ATAAAAAGAT	TGAAATATCT	720
TTATTAGAAA	AGAATCTCGA	TAATTTAGAA	AACAGAATAA	AAATCATAAC	ТААТАААТАТ	780
AACGTTGAAA	AAAATATATT	САААСТТТАА	CTATACAATT	AATTACACCT	TAAAAATGCG	840
ТТАСАТАААА	ATTAAGGACT	АСТАТАААТА	GAAAACACCA	САТААССТАС	AGACTCTAAA	900
GGAATAATTA	AATCCTCATA	TTTCAGTTCT	CCAAAAGTTT	AAATAGGGGC	CTTTTACTTT	960

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CATTATTATT ATTTTATTCT TTATTATTAC AAGATAATTC AAGAATCTAG ATTACAAGAT	1080
ATCAATCCTG CCATTAGTAG TTCAATAAAA CATTTAGAAT ATTTATACAT TATTTAATGT	1140
ATTTTTTCA TTTTTGAAAT AATATTGTTA TAACTTAACT	1200
TCAACTTGAG AATCCGATGT ACATAGAATC TGAACATCTC CTCTGCCCCA TTTGCCAATA	1260
TTCTTAATAT ATCTAGTAAA ACCCTCTTTT AAAATTATTT GATCTAGAGC AACAGTAATA	1320
GTAATATTAA TTTTATTTAC CCCAGGTCTA AAGCTAAAAT CTACAAAATA TCCGCCCTGT	1380
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AAAATAAAAT CTTCTAATTC TTTATATATT GCTTTCATAT CGGAATTTAA TTTTTCAAAT	1500
TTTTTTAAAT TTTCGGTTTT AATATTATTA TCTTTTATAC CAGAATCTGT GTCATCTTCT	1560
ATGTCACTTT TCTTGCTGTT TACTAATACA TCGCTTTTTT TTTCATCAAA AAACATACTA	1620
AAAATATTTT TAATAATATC ATTAAATATT TTATCTGAAT ATGTTTTTTT AAAACCAATT	1680
TTAGCTTTAA AAAAATCAAG CAAATCAACA CTTGGATTTT TTGTTTCCTT TTTTAAATAA	1740
GCTGAAAATT TGTCTGTATA TTTTTTTTTTT AATGCAAAAG ATCTAGCCTC TTCAACATTC	1800
AAAGAATTTC TAGAAAACTT TTTAAGATAT TCAAAATCCT TAGATGTTAA TTTTTCTAAA	1860
TTAACAACCA TAAAAGGCTC ATTGTCTAAC AAATTATCTT TATCTAGGTC AGTATAGAAT	1920
CTATATTCTA TGCCATCTGT TAATATACCA AATTCAACTC TCTTTGCTTG AGAACGAATA	1980
TTTTCAAAAT AAGGTTTTAA TTGCTTTAGA TGATTTTCAA GCTTTTCCCT GCTATTATGA	2040
TATTTGGCCT CTATTAAAAT AGTGGGTTCT TCATCCTTTT TTGTTGGATA AATAACATAA	2100
TCAACCCTTT TTAGTCCATC TTTAAGAATA TCTGCCTTCT CTTCAACTTT AACAATTGAA	2160
ATATCAGTAT GATCATAGCC CATCGCATCT AAAAATGGAT CAATAAGATT TTGTCTTGTT	2220
TGTGCTTCAT TTTCAATAAG ATCCTTATCC TTTTGAATTT TTCTACTTAC AGCTTTTATT	2280
GAATTTTCAA AATTTATATC TTTGTATTCA TTTGGCATAA TTATATTTTA CCAATAAAAT	2340
TAAAAATTAA TAATTCTAAA AATAAATTTC CAAAATGTTG TCTATTTTAA ACTCTTAACT	2400
GATACCTTAA TTCTTTTTTC TACCTAATTT TTTAGTTTAA AATCTTATTT TTTAATTTTA	2460
TTATTTTTTC CTTACCTTAT TTATACTAAA ATTTTTAGTA TTTAGCGAAT AATTTTCATA	
TCCTTTTATT AAAGACAAAA TATGATTTTC TCTTTTTTGT TTTTTAATAC CTTAAAATCA	2580
CTAAGCAAAG TAATAAAGTC TTCTTTGGTT AATGAATAAA AGACTAGCTA TAATAAAATT	2640
ATTTTATTT TCTTTACTAA ATTCAAAATG CTCTAAATAA AGCAAATTAG AGAAATTCAA	2700

		159			
AGGATCATTT TTAGCTATT					2760
AGCCAAAATT TCTTCTTTC					2820
GTCGGTATTT TAAAACAAA					2880
TATAATAAAC AATTTTTTA	T AAAAAGATA	T TGGTATTTT	C TCACAATTC	A TATCTATTTT	2940
ATAGAAACAC AATAATAAT	T TTTAGGAGA	r aaagtgcta <i>i</i>	A TCATGGTTC	T TTCATTTGTA	3000
TTGCTTGCAA TTCTTCTAT	A AAATATTCT	r TCATTTGGGT	C ACTGATCATO	TTTAGTTAAG	3060
ATTTTTTCTA AATCTTCTT	T ATATCCTATO	CATAAAAGCI	TATAACCTTC	TTTTACATAA	3120
TCATAAGTAA AAAATCTTA	A ATTAAATTG	A TAGATATTAG	CCCCAGAATA	AAGAAATATA	3180
AAGTTTTCAT TATTATATT	C CTTTAATAA	A GATTTGCGAT	TCTTTATACI	TGGATCTGGC	3240
ССТТТТТТАА ААТТААТАТ	C TTCTTTACTA	AGAATACTAA	ATGAACTAAA	TATTTTGTTT	3300
AATTTGGCCC ATGTTTAAT	T CAATTCCTT1	ATAAGGATTI	TCTTTGCAGT	CTTTTAAGTC	3360
TCTAGTTATT CCTTAATAA	T ATTATCACTA	CTTTGAATAA	CAAATTTTGC	ТТТААААТТТ	3420
AATGTAAAAG TTTATTACT	A CGAGGAAATA	TCGCAAATTT	AAAACTTGAA	TGCATATCTT	3480
AAAACCTTTT TTTGTTTTC	A AACTGATAAA	TAAGTTAAGT	ТТАТААТТАС	ТАААТАТАТС	3540
CTTTCTTAGC AAGCTAAGA	C CAAATATCAC	AATAGAAGTA	ATTCTCAATA	AACAAAATAC	3600
AAAAAGTAGT TATCATATCO	G TCTTTAACCT	TAAATAAGGT	TGCTATAAAC	AACCAAGATA	3660
TTTAATTTCT TTTAAAACCC	TTATTCAATC	TTTTTAAGCA	TAGGATCTTA	TAATTATAAG	3720
AATATAATTT TATTTACATO	С ТСТАТАТТАА	TAGAAAGATG	CAAATATGTG	ATCAAATTGT	3780
TATTTTTGTA ATATGGAATA	GTCCTTTATA	GGGACGCTTA	ATGCTCTATA	CTTAAGATTG	3840
GAATTCTCTA TGAAAATATA	A TACTCGCTAC	CCATGTAAAG	CTGACTTATT	TTAGCACGTA	3900
TCGCTTAAAC AATTATATTT	ATATTATCTT	TTATAAAGTT	AATTTTTTCT	TGTAGATTAT	3960
TTTTTAATAA AAAAGGCACA	AATTACCACA	ACAAGTTCCA	GTATAAATTA	ATAGTTCTTA	4020
TCTCAACACT AAAGTACATA	AACATCAAAT	АТСАААААТА	TATAAGAACA	ACATACTACA	4080
TTGTTTTAAT GAAAACCTTA	AAAGGAATGG	TTAAACTCTC	ATTAAGCTAA	AACCAATGCA	4140
ААААТАТСТТ ТАТАААТТАG	CAAAAGAACT	AAAAGTCACA	AACAACTACC	ATAAAAATTT	4200
GGTAGTAAAT TCTGGAACTG	AAATTTACTA	ТАААСТСААТ	TATTCTAAAA	AAAATATTGC	4260
CTTAAATTAA AGAATGCCTT	AAAAAAACAA	AATGCTCTGA	ТТТАААССТА	ТАСССААААТ	4320
ACAAATTTAC TAAAGAAGAA	GATATAGATT	TAGAGAAGAT	СТТААТААТА	AAAATATTAA	4380
TATAAAAGTT GCTCAGTATG	CTAAAGGCAA	AGAGTTTAAG	TCAAGTTTAG	AAATTACAAA	4440
GAGTAAAACT ATAAACTTCC	TTTAAGAATG	AAAATTTATT	ТТТАТАСТТА	CTTGGCTTAA	4500



TGTTTATAAA GAAATCTTAG AACTTGCAGA TTTAATACAA GCAGAGGTGC ATGTTGCAGG 6300 AAGGATAAAT AGCTATATAA AAAAAAGAAA GACCACTAAA GAAAAAGAAT ATAAGAAGAG 6360 AGAAATTAAG AATAAGATAG AAAAACAGGC TCTAATTAAG TTGTTCAATC AGTTATTAGA 6420 AAAAAGAGGC GATATTGAAA ATCTTCATAC TCAATTAAAT AGTGGACTTA GCGAGAGAGC 6480 ATCTGCAAAA TACTTTTTG AGAAAGCCAA AGAAACTTTA AAAGCTGCTA TTACTGAAAG 6540 ATTAAATAAC AAACGTAAAA ATCGGCCATG GTGGGCAAGA AGAACACATA GTAATTTAGC 6600 AATACAGGCA AAAAATGAGG CAGAGGATGC TTTAAACCAA TTAAGTACTT CTTCTTTTAG 6660 GATACTTGAA GCAATGAAAA TAAAGGAAGA TGTAAAACAG CTTCTTGAAG AAGTAAAATC 6720 TTTTCTAGAT TCTTCAAAGA GCAAAATCTT TTCTAGTGGC GATAGATTAT ATGATTTTTT 6780 AGAGACGAGT AAATAAAAAA ATATATTTTA AAGGCTAATA ACTTAAAATC AAAGTCTTCT 6840 GTTAAAGGAA GACTTTTTTA TAATTTTATT TAAATAACGA AAAGCTTGAT AGTTAAAAAA 6900 TCTTTTTTAT TAAAAATATG TTTACTAAAC AGAGCTCAAA AATGACTATA TTTAGTATCT 6960 CTATAAAAGA ATTTTTCAAT ATTTTAAAAA ATTTATAGAT AAACATAATC TAAAACCATG 7020 CATTAATACA AACCTAAAAC ATACTTGGTC ACTTGTAAAA GTAAATTGTA TCTAACTTTT 7080 TTTATTTATT GAATATACGT AAAAATTCTT TATAATTTCT ATTTTAAAAC GCTGCTATTT 7140 7200 TTACCCTCTG TTCTAATCCT ATCAAACAAG GTAATAAATT CTTTAAATTT CTAAAAAGCCT 7260 AAACTTTAAA AGAACTTGTC GAAAATAATA TTTCTCTTAA AAAAGGTTCT AATCTTTTAT 7320 TTATAAGAAC TTTTATACTA TTATAAAAAT GTATCTTGCC TTGATATATT TGTATTCTTT 7380 ATAAATCAAG CCTTCTACTT TTTTTAAGAA TATTTCTATT TTTTATAAAC TAGTTTTCTA 7440 CAATAGAAAA GAAATAACCC AAAGCCCTAA AAACTTAAAT AAATGTTAGC TATAATAACT 7500 AAAATAGAGA TAAAAAACTC AATCATAAAT AATGGTAAAA CAAACTTAAA CCACGTACCA 7560 TAACTCAATC TGGATATCCC CAATACAGCC ATTATAACTC CGCTGGTAGG TGTTATCAAA 7620 TTAATAAGCC CAGATGCAGT CTGCATGGCA ATAACAACTG AAGCTCTTGG AATTGACAAA 7680 AAATCGGCAA GAGGAGCCAT TATTGGCATA GTGAGACTAG CATGTCCTGA TGAAGATGGA 7740 ACAACAAATC CTATAAATAT TTGAATAATT TCATTCAATA TGATAAAAAG GGGTCTTGGA 7800 AGATTGTATA AAAAATTAGT AGCAGCATTT AACATAGTAT CTGTAATCAA CCCATCATCA 7860 CATACTATCA TAACACCTCT AGCAAGTCCA ATAACAAGAG CAGCGGTTAG CAGACTTTCA 7920 GAACCTTTCA CAAACGCATC CCACATTTCA GTTTCACCTA ATTTACAAAT AAAAGCCGAT 7980 ATAATAGCAA CTCCAAGATA CAACATTGTC ATTTCTTGCA TCCACCAACC AAGATTAACA 8040



እርጥልጥል ጥጥ ጥጥ ጥልርርመመጠልል	T ITTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	103			
ACTATATTT TACCTTTAA					9840
TTAATCGATA AATCAAGTA					9900
AAGCTCCTAT TTCGTAAAT	T ATACATCAA	CACCAACCT	TACAATCAA	AAAAATTTTT 3	9960
CTCATTTAAC TCATGCTTA	ACATGCTTA	TAAATTAAA A	A TCCTCTCTTA	CTAAAGACAT	10020
AGACATGCAT CTTGGCCCAC	CACGACCCC	TGAAAGCTCC	S CTAGACGGAA	TTCTGTGAAC	10080
TTTAATACCA TTTTCTTCAA	ACAGCTTATT	r agttacatg <i>i</i>	TTTCTAGAAT	' AAGCAATTAC	10140
TTCTCCTGGA GCTATCGCCA	AAACATTAGO	ACCATCATTC	CATTGTTCTC	TTGCACCATG	10200
TATTAAATCT CCACCCGCAC	ATTTTATTA1	GTCAATTTTI	CTGCCTAAAT	AAAAGCTCAA	10260
AACATCTTTA AGCTTGGCTT	TTTCTTTTT	T AATATTAATI	TTATTAGAAT	TTGAATTGTA	10320
AGTTAAAACA TAAATTGAGA	AATACATATO	ATCACTTGTA	AAACTTGTAA	AAACGCTATA	10380
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CTCATCCCCA CCTTCCAAAG	AAGTTTCTTC	ССАТСТАТТА	AACCAAATTG	GAACATTTTC	10620
TTTGTAAGCG GAATGATATT	ТАААААТАТА	CTCTGCAAAT	ATTGTCTCTC	TACGTCTAAC	10680
CTTGGTATAC ATTTTATTTA	TTGTAATTCC	ATTGCCAATA	CTGGCAAAAG	GATCTCTGGT	10740
AAATAAAACA TTGGGCATAG	GATCAATAAC	AAAAAGACTT	GAACCATTAA	CCCAATCATC	10800
AAGCGAAAAT TCACAATCTT	TAAGCTCTTC	TCTTGCAACG	CCGGAAATCA	TTTTAGAAAC	10860
CATATTATCA ACGGTTAAAT	TAGAAAAATA	ATCTTTTAAA	АТАТТААТТА	CACCATCTGT	10920
TTTTATTTCT GCTTCCAGAA	TAAATTGAGA	ТАТАААТТТА	TTTTTGAGCG	CTACAGAAGA	10980
AGCAAGAACT TCACTAACAA	GATCCTCAAC	ATACTCAATT	TCAACTGAAT	TATCTTTTAA	11040
AATATTTACA AAAACTTCAT	GCTCTTGTCT	TGCAACTTTA	AGATAAGGAA	TATCATCAAA	11100
ТАААААТТТ ТТСАТААТСА	AGGGTGTCAA	ATTTTCTAAT	TCTTCTCCTG	GCCTATGAAG	11160
CAAAACTTTT TTCAAACGAC	CTATTTCCGA	AAATATATTT	ATTGGATTTA	AATATTCTTC	11220
TTCCATCGAT TTCCCCCTTT	ATGAAAATTG	ТСАТАТАТТА	AAATACTATA	GTTTATATTA	11280
AAAAACATCA ACTATTTTTA	ATAATATTAA	АААТАТААТА	ТАААТАТААА	AAATTGAAAA	11340
AATAAAAGTT CTAAAAAACT					11400
AATCATGAAG AATATTAATA					11460
CTCTTGTGCC TTAATTGCAG					11520
AACACAAAAA ACACTACTAG					11580
					11300



	165		
	CAAAAATTTT CAAGGAATTA GCAATAATTT		13380
CTTGCCCTAT ATACAGCTC	СААААТАССА ТТААААСААА АТАТААТААА	CATATTTGGG	13440
GACAGTAAAT TAATAATTG	CTATTGGTCA AAAGGAATCT ATAATAGCAA	AAAATTAACA	13500
CAAATTACTA TTAATTTAA	CAAAAAGACA ACTGAACTAA GGAAAAAATT	TGAAGAACAA	13560
GGTGGAAAAA TTTCTTTTA	TCCAGGAAAT GAAAATATTG CAGATCTTGG	ТТТТСАТААА	13620
ACTAAGTAGA AATATTGTC	AAAAATACAT AAAAACAATA TTTCTGATTT	CAATGGTTTA	13680
TTTTTATTGT TGTACGACA	TAAAAATAAA CCATGATTAT GAAACTGATT	TTAAAGTTCT	13740
AGAATCTCCC TCTAAATACA	TCAATATAGA TGTAATTAAA GCTACAAATG	AATATATTTA	13800
TATTCAAATT ACAAACAATA	GCTTAGACGT AGTAAAAATA AATTGGCAAA	ACACTAGTCT	13860
TAACAACGAT AAGATCGTCT	TAAAAAAAGA AGATCTTACA ATAAACAATG	AAACAGGGTA	13920
TAAAAATAAA TACAGAGAGI	TTTTTATTGG TCCTAAAACT TCATTTAAAT	TTAAAGTATA	13980
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TCCGTCTATT TTTAAGCTCA	ACATAACAAA AGTAGGAATT GAAGCAAAAA	AAACAATAAA	14100
TGTTTTAATA ACAAGAACTA	CAAAAATTAA TATTACTAAT AAATGAAAAT	САТТАТТАТТ	14160
TTTTTGTTTT TCTATTAATA	ATAAACTCGT AGTTTGTATC TTTGTTGTTT	TCAAATGACG	14220
CTTTTATAGG AGCAAAAAG	TTCCAAGAAA TATAAATTGT AAAATTATTT	СТССАААТАС	14280
GAGAATAAAT CCCATCAGAT	CCCCTTAATA AATCTGGTTC TAAATTATAT	TCTCCAACAA	14340
ATTTCCAATC ATAAAATTG	AATTTAAAGC CTGATGAAAA TTTTTTAATT	TTAAAAAGTG	14400
AATCTTTTCT GTCTTGAGAA	TTAAAGAAAT TGAAAGATTT TGATAAATCA	ACAAAGAAAT	14460
TAACAGGTTC TAGACCAATT	TGGTCCATAT ACCCTTTAAA ATATTTAAAA	GTCTTAGTAT	14520
TAATAGATAA AGTAGAAAAG	TAAATTTCTA AAAATTCTGT ATATTTAAAC	TTCAAAGTCA	14580
ATGCAGATCG AAGTTCATTA	TCCGTAAATT TCTGCAAATT TATTTTCCAA	CCAACATCTA	14640
CCCCCAAGGT AAAAGAAAGC	TTATTGTCAA AAAAAGTTAA AACGTACAAT	PCCTTTTTGT	14700
AACTAGAATC TAAAGAATAT	GGAACAAGTT TGGTTGTAGT ACCAATCTTG	GAAAAATCTC	14760
CTTTTAAAGG ATCATAATTA	TATTCAAAGT CGTCTTTCAT AGCAAACAAA	AATTGAAAAT	14820
CAAAAACATT AAGCTTAAAA	GAAAGTTCAG AAACTCTATT TATCAAAGGA	PCATAGGCGA	14880
СТААААААСТ АААТТТАААА	TAATCCAAAT ATCTCGGCTC AATTTTATAA	FACAAAGCAG	14940
GAGACATTTC TAAATTTTTA	TAAGGCGATG ATGGTTTTTG AGGCTCCAAA (GACTTTGAA	15000
CAGCAGAAAT TCCAGAGTTT	TTCATAGCAT CTTCTTTAAA CTTTTTATAA 1	CATTTAATTC	15060
CAATCCCAGC TTCTTGTAGC	AAATAAGGAA AATCTAAAGA AAGTTTAAGC 1	CAGAAGAAG	15120

CTTTAATATC TTCAAAACTA TTTTTTAATT CACCTGAAAG CTCAGTAGTA AAATAATCAT	15180
AATCATAAAT TAAAGAGGCT GTTAAACTTT GATAAAAAGT TTCCGGATCA GATAAAAAAA	15240
TACTACTATT CTTATTAACC AAAGATTTTA CATCAGAATC ATATTTTTTA TTAAATGAAT	15300
ATAAAGTAGC CTTATTTTCA AACTTTAAAG TACTTCTAGA AAATAAAGGA TATCTAATAA	15360
AAGGAAGCAA GTTTAAATTT ATTTGGTTAA TAATAGAGTG CTCACTTTTT TTATCTTTAT	15420
CTTCAACTTT AAAATCTTTA TTTAAAGGAC TATACTCAAT AGTATTAAGA TATAATAAAT	15480
TTTCAAAAGT AATTAAACGA TTGTAAAAAT CAGCATGAAT TTTTATATCC GTTTTATTTT	15540
TTATATCAAA TAAATAATTT TTTATTTCAT AATTAAAGTC CTTTGGACTT GTTATGCCAT	15600
AATTATCAAA AAAAACATTA TTTCTTAAAT AAGGATTAAT GCCAAACCTA ATAAAAAAAG	15660
AATCGGATTG ATCAATATTT TTTAAAGTAA TTGGTTCTGG AGGAATATAT AAATCTTTGG	15720
TTAATTCTGT TGTTTTTTTA GTATTTTTCT CCTTCACACT CTTTTTATCA TTATCTTTAT	15780
CTTCTAGATT TTTAATTTCT GGGCGCATTA TCATTTCTTT AGTATCAGCT GGAAATGTCC	15840
ATTGGTTATT GTAAAGATCT TTTTGAAAAT TCAAATCAAT ATATGGAGCA TAAATTCTCT	15900
CCAAATAAAA CCATTTTCTT GTAGGATCAT TAACATCTTT TGGTTTCTCT AAAGGAGATT	15960
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AATCTAAAAT CGAACCGTCA TTAAATGTTC GCTTATAAAA AGAAGATAAA TTCCAATCAA	16080
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CTAAAACAAC CGAGAAAAGT GCATCACTTA AAAGAAATTC TGTTTTAAAT TTAAATAAAT	16260
ATCTAAAAGG AACTTCAAAC CCAAATACAT CTCCTTTGTT AAGATTGGAA AAACTAAAAA	16320
GAGATTGTTT TAAAGTCCTA TTATCAAAAG GATAATATCC TCCATCGTAA CTATAAACAT	16380
TCCTGGTAAA ACCCAATCCA AAATTTCCTT CCAAAGTTTT AAAATGCCCC AAAGTATTGC	16440
CCAAATTAAA ATCAATTCCA GAATAAAATC CCAGATTAGC ATAAATGTCA AAAATAAGCT	16500
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AAGAAGAATC TTCACTTGAA GATTTATTAC CAAAAAGATA AACGGTATTA AAAACAGAAA	16680
AACCTTTTCG TGGATTTAGA CCTAAAGATG GATTAAAAAA CAAACTATCT CCCGGTCTGA	16740
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CAAAATCTCC CGAGGGCAAT GCCCATATTT TAGAAGCCTT GATTGAATAG TAAGGCTCTG	16860

		167			
GAATTTTACT AGTTGTTG					16920
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CATAAAAATA AAGCTTTT	CA TTGGTATCCA	TATCAAGAAT	ATATTCAACA	TTTCCAATAG	17100
CATAAAGTTT TTTAGAGT	TC TTATTAAGGA	CTATTCTGTC	GCCTTTAATA	TTGTGCTTTT	17160
TATTTTCTTT AATATCTTC	CA ACCAAGATAT	' TAACTCTTCC	ТТСАААААТА	ATACTTTCAT	17220
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TGCCAACCCA CCATCAGGA	С ССАААТСТАТ	ТАТАТААТСТ	GCCTGTTTAA	ТТАСАТССАА	17700
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TGATAAAGTT GTTGCAGAT	T GTCCTAATTT	AATATATTCA	AGTCCAACTT	СААТТААААА	17940
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CATCTCTAAA ACATCATGT	A TATTTTTCC	TTTGTATCTA	ACTTCTAAAG	ТТТСТТСАТТ	18060
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GCCAGAGCCA CTCTTGCCAG	ATATTACAAC	TAAACCATCT	TTTGGAATAT	CTACATCAAC	20340
ATTTTTTAAA TTATGTTCTT	TTGCTCCTCT	GACAATAATT	ТТТТТТТТСА	AACTTTTTTC	20400

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СААААТТА	C ACCTCTCTT	TTTTATTAC(G AGCTATACTA	ATTTTGCTAC	TAAGCTCTTT	20460
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GCTTTCAAGG	TACACACAGC	TGATTATGTT	ACAACTGACG	ATGGAACAGG	AATTGTTCAT	27480

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ATTGCTCCTT TTGGAGAA					
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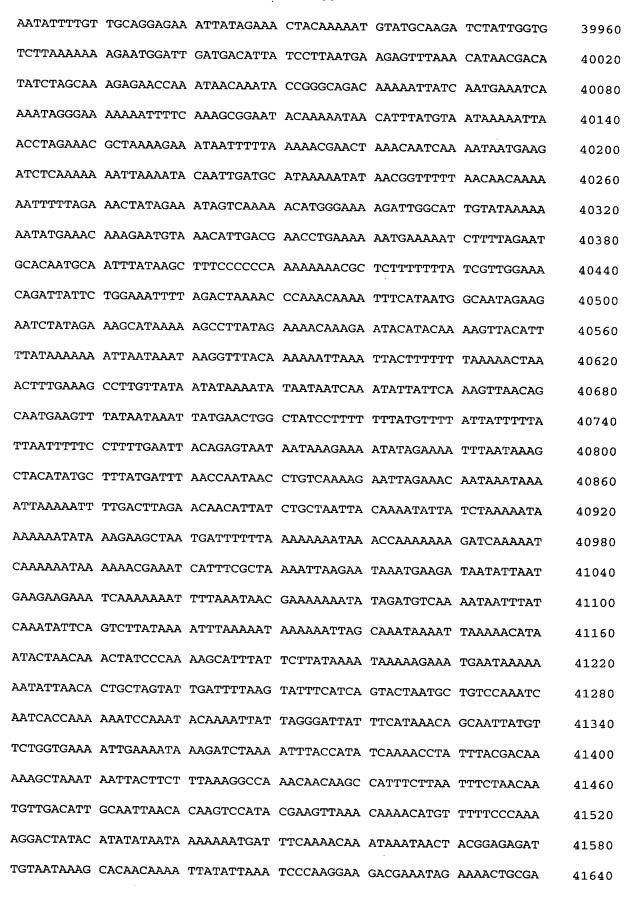
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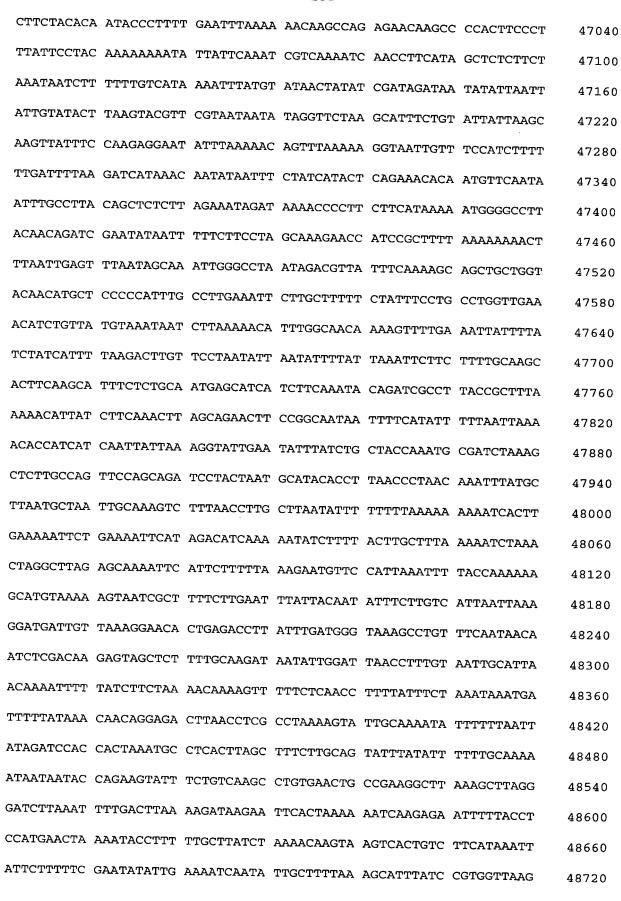
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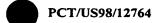
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ATATATCGGC A	GCACTTAGA	ACTTCAACAA	AATCGGCAAA	AAATTCTTTT	GTTCTTGTAA	45960
AGGTATGAGG C	ATAAAATCC	ААААТТАТАС	GTTTATTCTT	ATAAAAATTT	TTAATACCAA	46020
AAAGAGTATT T	PTAATTTCC	CTAGGATGAT	GAGCATAATC	GTCCATGTAA	ATCACTCCAT	46080
TTTCCTCTTT A	ACAACTTCA .	ACCCTTCTTT	TTATACCGCT	ATAATTTTTT	GCAATTCTCT	46140
TTATTGCTTC T	ГСААААТСА	AAAATTGATT	TCCCATTACT	TTCTAAGAAA	AGATTTAAAG	46200
CCAAAAGCGC TO	GCTGAAAAA '	ТТТААТАСАТ	TATGAAATAA	AACAGTCTTA	AGCTCAACAT	46260
TTAACAAGCC TA	AAAAAAGAA	AAACAAAAT	ATTCACTCCT	AACTGCAATA	TTACTTATTT	46320
GAAAATCAGA TA	AATCTCCA	GACCCATAGC	ТАААААТАСТ	TATATCTTTT	CTGTTGATTT	46380
GCCTTTTAAT TI	TTAAGCAAA '	TTATTATCAT	CGGAATTAAT	TATCAATATT	CCATTTTTCT	46440
TTAAATTATT AA	ATATACTGT A	AAAAAAGCCT	CTTCAAGAGC	CTCATAATTT	TAAAAAATT	46500
CAACATGCTC GT	CAGTCAACA	TTGGTTAAAA	TAAGCATATT	AGGGCTAAAA	ТТСААААААТ	46560
GTTTCTTATA TT	CACAAGTT !	TCAACAATAA	AAATATTGCT	AATACCTGCT	ATTGCAGAAT	46620
TATCTTTAAA AT	CTTTAACA (CTTGACCCCA	СААТААСАТТ	GGGATTTAAT	ССТААТТТАТ	46680
TAAAAAGAAC AC	CTAAAAAC (GCCGTAGTGG	TAGTTTTACC	ATGAGAACCT	GCAATTCCAA	46740
TGCTATAGTA CT	TTCTAGAA 1	AGCTCTCCAA	GAGCCTCAGG	ATAAGATAAA	ATAGGTATAT	46800
TTAATTCTTT TG	CCTCAAGT A	AAAACTTGCA	AACCATCCTT	ATTATAGGCT	GAAGAATATA	46860
СТАТТАААТС АА	AAGACCTA 1	TCAAGCTGTT	TTAATGAAAA	СТСАТАААТА	ТТАТСАТААТ	46920
AAGATATTTT AT	TATTACTT A	AAATTTCAT	CGGTATAAAA	TTTATCAGAA	ACATCTACCC	46980



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TTCAAATTTT	GGGCTAAAGA	TTTGTACAAA	TAAATTTTAT	' CACCTTTGCA	AACTCTGCAT	48780
					TTTTATTATA	48840
CTCGCTTTAG	AAAAATTTAA	AATTTTAATT	AAAATTGAAT	CTAGTCGCTT	GCCATTATCA	48900
TTAGCAAGCA	СТТСТАААА	AATATATTTA	TCCAAACGCA	AAAAATACCC	CTAACAAACC	48960
TTACTATTTT	TTTTACAAAA	AAAATTAACT	' АСТАААААТС	TAAATATAGA	AACAAAAAAT	49020
GATGGAAAAA	CGGGGTGAAA	AAACCAAATA	TTTAAACCAA	AGAATAAAAT	GGACAAATAA	49080
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AGTCCAAAAA	CAATAATAGG	GAAAAACGAA	ACTTCCAAAG	CTCCAAAGGC	AAAAATATTA	49200
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ATAAATATTG	ACTTTATTAA	AACAGATGTT	ATTAATAGCA	AATTTGAATC	CACTGTAGAC	49380
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AAAGCCACAT	TTAAAACAAC	TTTATCATTT	GGACTTAAAT	CTGGAAAAAG	AATAATAGCA	49500
АААААСССТА	TTAAATGCAT	CAAAACAATT	AAAAAGCTAA	TAATAAAAGT	AGAAATGGGA	49560
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TGCCCTAGTA	TTCCTATTCC	ТАТТААТАТС	CAAAAAGAAA	TTATATATTG	TGGCTTTAAG	49680
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TTATTAATAC	CCCCTCCCAA	ATCTAGCATC	TTGGAAAACA	AAATAACGGA	TGAAACTAGC	49800
ATTAAAAATC	CTTGAATCAA	ATCCGTATAA	GCTACTGCCT	TAAAGCCGCC	ААААААТАСА	49860
ТАААТАААА	CCAAGAAGGC	AAAAAAAGTA	AGACCAACTA	CGTAATCAAT	ACCCCAAAAA	49920
ACTTCTATAA	GTTTGGCACC	ACCTATTAAT	TGGGCAGAAA	TCAAAAACAT	TGAAAAAAA	49980
ATCAATACAA	ATCCACTCAT	TAACGCCAAA	AAATCACTTT	CATATCTATG	ССТААТАТАА	50040
TCAATAATAT	TAATTGCATT	AATTTTTTT	GATTCGCGAT	TTAATCTCTG	ACCAACAATA	50100
ATAAAAACAA	TTAAAGTTGT	AGGAATTTGT	ATGGTAGCTA	АТААТАТААА	AGATAATCCA	50160
TACTTATAAA	CAGCAGAGGG	ACCGGAAATA	AAACTACTAG	САСТААТАТА	GCTAGAAGAA	50220
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AACAAAAACC	TACCTCTATT	TCTTTTTTA	AGAAAATCTA	ААААТАААА	ААТАТСААТА	50340
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GGAATAAAAT '	TCCTGACAAA	AAAAACCACA	AAGGAATATT	AAATATAGTA	GTTGATGTGT	50460
CAATAAAATA (GGCAAAACAA	AACCACAATA	САААСАТААА	AACATACAAT	AATATAGCGT	50520



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TCCTTTGCAG	AAGAGTCCCT	ATGTCCTTTC	ATTGCATTAA	TAGCTTTAAA	CCTAATATTA	52320
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GCCTTAATTG	ATGGGCCCTC	GTAATTATCT	AGCGAAATTT	САТАААТТСТ	ATCCTGATAA	52560
TCAACAGCGG	ACATTTTTCC	AAGAGCAATA	AGTATTTCTC	TTCTAGCCCC	ATCATTTCCA	52620
GAATATTTTT	CAAAAACTTC	CATCATGTTT	TTAGAATACT	CAAGAGAATT	AAGCTCTCCT	52680
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CCCCCGAAT	ACTTAAGAGA	ААТАААСААТ	TCAAGTATTT	CCCTTTTAAG	CTCAGCATTA	52920
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ACAGGCTTAT	TTTCTGTAAT	TTCGGGCAAC	AAAGGCGGAC	TAGGAAGAGC	TGGAGAATTA	53160
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GGAAAAAACC	AAACATTGAA	TAAAAAAGCA	GACCTAAAAG	AAAACATTTC	AAAACACTCG	53700
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ATGAAATCTG	CTGAGTAATA	ATTGAAATGG	GAAAATCCAA	GAATCATACG	AAGCCATCTG	54060



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GTATCATCCT	TATACTCCTT	TCTACCAATG	CACATAATCC	CATTATCTGT	CAAAAGAGAC	55020
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GCCCCAAAG	TACCAACTGG	CATAAAACAA	GGAATATCTA	CTCTACCATG	AGGAAGATTT	55800

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GAAGAATCAA	TCTAAATGAA	AAGAATATCC	ТАТТТААТАА	АТСААААААТ	GCATCAGAAT	56160
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CATCAAGAAC	ATCGTAGGCG	TTCTCCTTAA	TTTTTTTACC	AACCTTAACA	AACTCTCTAA	56340
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CCCCAGATCT	ACTGTCGCCA	ATGCTATTCT	TAATAAGAAC	ATCTCTTCTT	GCTATAGTAT	56580
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TATTCAAATA	ATAATTTGTC	AAGCCAATA	TTTTTTTAGO	ATAACTTTCC	CTGGCTTTTT	59460
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TGAAAGCTTT	GTGTATTTTC	CCACAAGACA	AACCAAAATT	CCACTTGCAA	ATTTATCAAG	63120
TGTGCCAGCA	TGCCCAACAC	GATTTGTATT	AAAATATTTT	TTTATAGGGA	AAAGAGTTTC	63180
AAAAGAAGTT	TTACCTTGTT	CTTTATTAAT	TAAAAGGAAT	CCATTTTCCA	AATTTAATTC	63240
TCTCTTGTAG	TATTTAATCC	TTCAATTAAC	TTATTAACAT	AAAATGATTT	GGAAAGAGAA	63300
ТСАТССТТАА	САААТААТАА	TTTGGGAGTG	CTTCTAACTT	TAATTCGCTT	AATAATTTGA	63360
CTTTGAATAA	ATCCCTTAGC	ATTATTTAAA	GCTTTAACTG	CATTGTCCAA	AGAAGCACCT	63420
TCCTTAATAG	AGCCCATAAA	CACTTTAGCA	ТТТАТТАААТ	CTTTTGAAAA	ТТСТАСТТТА	63480
ACCACGGTTA	AAAATGAATG	AATTCTGGGA	TCTTTAATCC	CCCCACTTAC	TATTAAATTG	63540
CCGATTTCTT	GAGCAATAAA	ACTTTCAAGT	TTAAACTTTT	TAATATTCTT	ATACATAAAC	63600
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CTTCAATTAT	ATCTCCTTCT	TTAATATTAG	CATAATTATC	AATCATAATA	CCACACTCAT	63720
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CAGCAAAGCC	AATAAACTGC	TGCTCAACAT	CTGGCTCAAG	CATTCCTTCA	AGAACTGACC	63960
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CCCCGGCTTG	AGGCATTGAA	GAAAATCCTA	AAACACTAAT	GGCTTTAGCG	GGTCCAACGC	64440
TCTTAACAGA	AACACCCTTT	TCGCTAATTA	ATGCCTTAAC	TTTACCATAG	CACGCTCCAC	64500
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CGCGCCCCAA	ATCAATCTTG	GCATCAAGCA	CTTTTCCAAT	AGCTCTTTTG	GATGGATTTG	64620
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PCT/US98/12764 203 GGCTCTAAAT TCACGCGCTT TTTCACTATC TAAAGAATTT ACAAAATTAG AATTTTCAAC 80640 ATCAGTGCTT AGTGCTACAA TTTCATTTTC AAGAAATTTT TTATCTCCTT GTGAATATAG 80700 CAAAGCAAAT GGCTTTAAAA AAGAAAAAAA ATATTCCTTA CCTATAAGAT AATGGTGCAT 80760 TGAACATCTT GTAGAGTAAT TTGGGTAAAT AGCAGATTCA ATCTCTTCAT CGCCACCAAC 80820 80880 AATAAGCAAA GGATCAAAAT AAGGAGCTGG AATTTCTTTA TTGTTGTAAA AATAATACCT ATACGAAAAA AAAGATTCTT CCATGCAATC AATGTGCCCA CAATAAGCTC CAAGCTTATA 80940 AATCTTATCT GAATATTCTA CTAAAAGTTT GGCAGTATCA TTGAAAGATT CTTTTCTAAA 81000 81060 ATTATAAAGC AGTGTAGGTA ATATAAATAA AATATTCTCA TTTGATGAAA TTTTATTTAA AACAAAATT AAACGTTCAT CATAAAAACA TGCCTCATCA ATAATAAAAG TGCCACAACT 81120 AGGATTAGAG GCTATTAAAT TTTCAATATC AAAAGAGTTG CTAGCATAAC CAATCTCATC 81180 AATTTTATCT TTTCCACCGC CTCTATATGG TATTACGTTT TCTGGATAAT CTTGAAACCT 81240 CCTCTTGTCG AGAAAATTTC TAATAAAAA TACATTAACC CTACTTCTAT TTCCTTTAAT 81300 AATATTGCCC AATACCTTGA AAGATTTTTT TCTTACAACA AGCGAATCTT TATAAATTTT 81360 TGCAGCATAT TCTGTTTTTC CACTTCCCAT GGGTCCAACT ACAAGAATTA AGTTTATTTT 81420 TACCCTAAAA TCAAAATGAC TAACAGAGAC AATGTTATTT AATTTAGTAT CTTCTTTATT 81480 AGCAAAGTCT AAACAAAAAC CCAAAAATCC TCCCTAAAGT AAATCAAATT CAATTATATA 81540 AATAAAACA ACAAAAAACA TTAACATTAA AAGCCTAAAA ATTAATAATT TAGGATCTTA 81600 TTAAAGCTAT TATTCAAAAG AATAATAGCT TTCAAAACTA TCATCATCTA ACAAAGCTTT 81660 CTTTATTTTT AGTTTATTCT TCTCATAAAT TTCAATATAA TTAAATTTTT TAGAACTATT 81720 81780 AACTTTTTTA TAAATTGAAA CCAAACTTCC AACCACATAA TTTTTAATTT CTACCAAATT 81840 AGATCCAAAA TATTTTTTT TACTTATCAA GCTTGATCTA ATATCCAAAA CAGAAGTTTT 81900 TTCAAAAAAT AAATTTAAAG CATTTGGAAT AAAAAAATCC CTAATATCAA TCAAGAACAA CTCTCTGATC CCAAAACTAT TAAAATTGCA GCCCAAATTG TTAAAAAGAT TAAACTTAAA 81960 AGCATGTAAA GCAAAGGTAT AAACATCTCT TTTATCAGGA TATTGAACCT CAATCATATC 82020 AACATAAGGA TAAACAGAAT AATTTATTTT ATAACCAAGC AAACTATTCT CATAATAAAC 82080 TGGAGACTTA TCTTTAAAAT AAATTTTATT AAAATATTTG CTATTTAAAC TAAATTTGCT 82140

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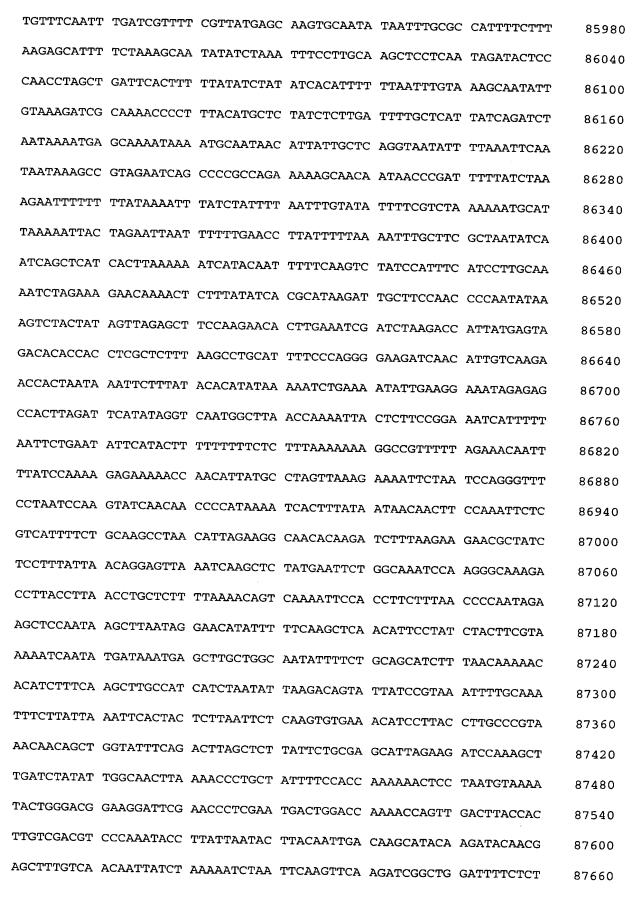
82200

82260

82320



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CCAACCCAAT ATCCTAAAAC	TTCATTTTCA	AAAACAGGGT	TTTCATCCTT	GCTAACAATG	93240
TCCTTTTCAT TTGAATCTTC	TGAATTTGCA	AATAAAAGCA	TTGAAAAACA	AAAAAAAAGA	93300
AAAAACTTTG AAAAAACTCT	AGTCATCAAT	CCTCCTTAAA	ACCAAATTAT	AGCTCTTTTT	93360
TTAAATTACT CATGTAAGGC	AGCTAATTTA	AAAATTAGCC	ACAAACATCA	TTATACAACT	93420
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AAACACAACA GGCCAATTAA	TTGTTATATG	GGACAATTAA	TGCTAATATT	AATAAAGTTA	93540
ATTGTCTTTA AGGATTTAAC	CGTGGTAAGA	GGAATTTATA	CAGCTGCCAG	CGGAATGATG	93600
GCAGAAAGGC GCAAGCTTGA	TACCGTGTCA	AATAATTTGG	CAAACATAGA	TCTTATTGGA	93660
TACAAAAAG ATTTGTCTAT	TCAAAAAGCA	TTTCCAGAAA	TGCTAATAAG	AAGACTAAAT	93720
GATGATGGTC TTTATAAATT	TCCCAAAGGA	CATCTTGAAA	CAGCTCCGGT	TGTGGGCAAA	93780
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ACTACTGGCA ATCCATTAGA	TTTAGCACTC	ACCGATCAAG	GATTTTTCGT	AATACAAACT	93900
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GTTACAAAAA GCGGATTTCC	CGTTCTAGGA	GAAAAAGGAT	ACATATATCT	TAAGAAAAAT	94020
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GATACAAAAA CATCTGGCAA	AGCACAAGAA	ATTGATATAT	CATTAAGGCC	TAAAATAGAA	94260
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CAAACCCATA ACAGAGCAGG	AACCCCTGCA	ACTGAAAATA	CTTTAAGACC	ACTTGGAAAT	94620
CAAGTTGGTC ACGGAACAAA	AATTGCTGCC	ACCCAGAGAA	TATTTGAACA	AGGAAAAATG	94680
CAATCCACAA ATTTACTCAC	TGACGTTGCC	ATTGAAGGAG	ATGGATTTTA	СААААТТСТТ	94740

CTACCTGATG GAACTTATGC ATATACTAGA GATGGGTCAT TTAAAATCGA TTCTAATCGA 94800 GAGCTTGTAA CAAGCCAAGG ATACAAAGTA TTGCCTAATA TACTCTTCCC AGAAGAATAT 94860 ATCCAAAACT CAATTACAAT ATCTGAAGAG GGAATAGTAT CGGTAAAAAT TGATACCAGC 94920 AACGAACCAA TAGAGCTTGG GCAAATTGAA ATATCAAGAT TTATCAATCC TGCAGGACTA 94980 AGTGCCATTG GAAGCAATTT ATTTAAAGAA ACAGCTGGAT CAGGCCAAGA AATAGCAGGA 95040 95100 TCTATTGCTG AAGAAATGGT AACAATGATA GTAGCTCAAA GGGCTTATGA AATAAACTCA 95160 AAAGCTATTC AAACTTCTGA CAATATGTTA GGAATTGCAA ATAACTTAAA AAGGCAATAA 95220 AATAAAAAA AGATTATTTA TTTTTATTTT ATTTTTCACA ACAAGCTCAA TTATAAGAGC 95280 TTCTCATGAT TTATGTTTCA ACATTGCGCC TAGTAAAACA TATTTCTTTT CAAAGAAGTA 95340 TTCAAAAATA TGTAACAATC AAAGCTTATC AAAAATATAT ATCCCCCCAC ATTTAACAAA 95400 AAAATCAATA ATTTTTGAAA TGATTTATTA CATTACAAAA AATTTATCAA ATGAAAATAT 95460 CTATATACTT CAATTTAACT TTGATGAATC TGAAATAAAC ATAGAAGATA AATTTTTCAA 95520 AAAAGTAAAA TTTAAGGTAA AAAGCAACAA TTCATACAAA AATATTCCAA TTGAAAAAAC 95580 TCTTGTTTAT TATGCAAAAA ACTTTGAAAG CTACAAAAGA CACAATTACA TCAATATGTA 95640 CATTGATGTA ATCGAGCCAA TTGTATTTGC AAAAGAAAAT CTAAAAAAAA ATGAAATCCT 95700 TAATGAGTAC AATACATACT TTAAATACAA AATTAACACA ACAAGAATAA ATGATGTTTT 95760 AAGTCTAAAT GAATTAAACA ATAGCAAATA CAAAGTTATA CGCAACACAA TCAAAAATGA 95820 AGAGATAAGA TTAAATAAGG TGCAAAAAGA ATAATACCTA ATTTTATCTT CCTTTTCTAA 95880 AAATTATTAT TTTAATCTCC CTTAATGCAG CTAATATTTA ACAAATCAAG GATTAATTAG 95940 TAATTTAACG AAAAAAGTTT CATTAATTGC AATAATTGAT ATAAAATAAT AGATATTAAA 96000 GAAATACAAT AAATAAGGTA AAGAATGAAC AAACTAATGT TGATGTTAAT TACATTTGCA 96060 ACGAGTCTAT TAGCCCAAAC AAACAAAGCT TCAACAGGAC TAAAAACAGA TCAATCATTT 96120 AACAATAGCC TATCTGAAAG CGTAAAATTA AAAGAAATTG CGGATATTTA TCCCACAAAT 96180 ACAAATTTT TAACAGGTAT TGGAATAGTA GCGGGACTTG CTGGAAAAGG AGACTCTATA 96240 AAACAAAAG ACCTTATAAT TAAAATTTTA GAAGAAAACA ATATAATAAA TGAAATAGGC 96300 TCTAATAACA TAGAAAGTAA AAATATTGCA CTAGTAAATG TCAGTCTCCA AGTAAAAGGT 96360 AATACAATCA AAGGTTCAAA ACATAAAGCT TGCGTTGCAT CAATACTGGA CTCAAAAGAT 96420 TTAACAAATG GAATACTTTT AAAAACAAAT CTTAAAAATA AAGAGGGGGA AATAATAGCA 96480 ATTGCATCAG GAATTACACA GCCCAATAAT AAATTAAAAG GATCTGGATA TACTATAGAT 96540

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CCCTCCTCTT	GACCTTTGCA	CAGACAATGG	AGCAATGATT	GCGGGACTTG	GATTTAATAT	100140
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215 GAAATCTACT TCATCCATCA ATCGGGGAAA AATTTAAATG ACCTAAGCGA AAAGAATTAC 101880 CTTAGAAGGC AATTTTTTAA CGCAGAAGAA ATGGCAAGTA TAGTTAAATT TTCTAATCTA 101940 ATAATAAGCA GAGCCGGAGC TGGAGCAATA AAGGAATTTG CAAATGCTGG TGCATGTGCA 102000 ATTTTGATTC CATTTAAAAA AGGCTCTAGA GGAGATCAAA TTAAAAATGC AAAATTACTA 102060 ACAAATCAAA ATGCCTGCAT TTATATAGAT GAAGATGAAA TTTTAAATAT AAATATTTTA 102120 AAAATTATAA AAAAAACTTT AAAAGATAGA GAAAAAATCA ACTCTCTCAA AGAAAATATC 102180 AAAAAATTCA ATAATAAGCA TTCTTCAACT TTAATAGCCA AATTGCTAAT AAAAGATATT 102240 AAGGAGACAA AATCTAAATG ATAATAAACG ATCCTGTAAA AATAACTGGA ATAGTAGACA 102300 TATTAATAAT AATAATTTTT ACATCTTTGG GATTTAGAGG ATTTTTAAGA GGATTTATTA 102360 AAGAAATTAG CGGATTTGCT GAAGTTTTTG TTTTAATCCT ACTGCTTTAC AAAAAAACTG 102420 AAGAATTTAG AAGGTTTGTT GAACCTATTA TTGAGCTATC CTACATTCAA GCACTACTTG 102480 TATTTTTTT GCTTATACAT ATAGGATTTT TAATACTACA ATCCCTAATA GAATCAATAA 102540 102600 TAAGTCAACT TAAATTGCTA TTCTTCAATA GAATACTAGG CTTAGTGCTT GGCCTACTTG AAGCTTTTGG AATAATTGCA ATCGTGGTTT ACATAATACA CTCACAACAA ATATTTAAAC 102660 CTGAATATTT CCTAAAAGAA AGCAAACTAC TTGATTATTT AAATCCTGGA ATAAACTATC 102720 TCTTTAAAAT TTCAAAAACA AAATAAGGGC CAGCAATGAC AATGCTTCCA AAAATTGCAA 102780 AAGAGATAAT AAACGAATAT GATCAAAAAA TACTGCCAAA TGCAATTCTT TTACTAGGAG 102840 AAAAATTTTC TTCAAAAAAG ATTAGCGCAA TTGAGCTTGC AAAAAAAATA TTAAACGGAA 102900 AAAACTTAAC AAACCCTAAT TTGCTCATTT TCTCAAATCT TGACACAGTA GAAGCAAAAG 102960 CACATCTTTC TACAAATTCG CAAAAGATAG CAAATAAATA CCTAGAATAT ATTAAAACTG 103020 TAATTTTTAC CAAATGTTAT TTCAGCAATG AAAAAAATTT AAAAAAAATA GAAAAAAATA 103080 103140 TCAACTACAT TAATTCTGTT TATTATGAAA AAGAATACAA TGAAAACATA AAAAATGAGC TTATAAAAA TATAGAAAAT ATAAACAAAG AATTAAATCA TAGCATTACT GTTTATGATG 103200 TAAAAAAAT TCAAACTTGG ATTTTTTCTG AAAAAGAAAA ACCAAAGGTA ATCTACATAA 103260 ACGAAATCGA AAATTTATCA TTTAATGTCC ATAACTCACT TTTAAAAAATA TTGGAAGAGC 103320 CTCCCTCAAA TATTTACTTT ATCTTGGCAG CAAGAAATAA AAACAAAATA CCAAAAACAA 103380 TACTTTCAAG ACTTAGAGTC TACAATTTCG CAAAACTAGA CAGAAGCTTA GAAATTCAAA 103440 GATTTAAAGA AAGCTTTCTA ATAAATAAAG ATATAACAAT TGAAGAGTAT TTCGCCTCAT 103500 TTTACAAAGA AGAAAGCAAA AAAATAAAAA AAGAATTGGC AAAAATTCTA AATATAATAA 103560 AAGAAAAAA ATCCATATTT AATCTTGAAG AAGTCGACTT TATAAAAGAT GAGCAAAGCT 103620

TTAAAATATT TTTAAACGAA CTTACAATTA ACATTAGAAA AGATTTTTTA GAAAACAAAA 103680 TAGATATTAA TCAATATCTA AAGTACACAG AGCATTTGAA AAATATTTAC AAATATCGCC 103740 CCTATAATCA AAATAAAAAA TTAATAATAG AAAACTTAAT GCTAAATTAT GAGGAGATAT 103800 GAATAATTTT TTCAAAAAAG CTTTAACAAA GCTAAACAAA TTATCTAACG AACAAAAAAC 103860 TAAATTTATT GAACAAATTT ACAAAAAAT AGAAATATAT GACGGAATAT TTGCATCAAT 103920 TAATGAAGGA ATCATTGTAC TTGACAAACA AAACAATATA ATCTATGCAA ACAAGATTTT 103980 ATACCAAATT TTAGCTTTAA CATCTAAATC AAAAATAGAA ATTCTTGATG ACATTCAAAT 104040 TCCAAACTTA ATAAATTTAA TAAAAGAACT AGTTAGAACA GAAGATAAAA TAATAGGATT 104100 AGAAGTTCCA ATCTCAAACG GCATATATAT TAAAATCTCA TTTATGCCTT ATGTAAAAGA 104160 AAAAAAACTT GAAGGCAACA TTATTTTAAT CGAAGACATT AAAGAGAAAA AAAAGAAAGA 104220 GGAACTATTT AGAAGAGTTG AGGCTTTGGC CTCTTTTACA AGGCATGCAA GAAATATTGC 104280 CCATGAAATC AAAAACCCAC TTGGAGCAAT CGATATAAAT TTACAACTGC TAAAAAAAGGA 104340 AATTGAAAAA CAAAAATGA AAAATGGTAA AGCTGAAAAT TATTTTAAAG TAATAAAAGA 104400 AGAAATAAAC AGAGTAGATA AAATAGTAAC AGAATTTTTA CTAACTGTCA GACCAATAAA 104460 AATTAACTTA CAAGAAAAAG ATATTAAACA AGTAATAGGC AGCGTATGTG AATTGTTAAA 104520 TCCTGGATTA GAAAATAAAC ACATAAAACT ATTGCTTAAT TTAAACAAAA TAAGCAATAT 104580 TCTCATTGAT GAAAAACTAT TAAAACAAGT TATTATAAAC ATCGTTAAAA ACGCAGAAGA 104640 AGCACTGCTT GAAACAAAAA AAGAAATAAA AAAAATAGAA ATTTTTCTCT TCGAAAAAGA 104700 104760 CAATAAAATA CATATCAACA TAAAAGATAA CGGAAACGGA ATAAAAGATG GGGTAAAAGA GGAAATATTT AAGCCTCAAT TTAGCACAAA AGAAAAAGGA AGTGGAATAG GACTTACTAT 104820 TTCTTATAAA ATAATAAAAG AGCTTGGAGG TGAAATTTTT GTGGAAAGCA AAGAGGGCAA 104880 AGGCACTATT TTTACAATTA CGCTGCCTAA ACTAAATAAA AAAAATATTT TAATTGAAGG 104940 GTATTGAAAA TGAGCAAAAT ACTTGTAGCT GATGATGAAA AGAATATTAG AGAAGGAATT 105000 GCTACTTATC TTGAGGATGA AGGATATTTT GTTTTCACTG CTAGTGACGG AGAAGAAGCT 105060 CTTGAAACAA TTGAAAATGA AAATCTTGAT GTAATAATAT CTGACCTGAG AATGCCCCAG 105120 ATATCTGGAG AAAAATTGCT CAAAATAGTT AAAGAAAAA ACTTGGGAAT ACCTTTTATT 105180 ATTCTAACAG CCCACGGAAC AGTTGATTCT GCTGTAGATG CCATGAGAGA GGGTGCTTAT 105240 GATTTTTTAA CAAAGCCCTT AGACCTTGAA AGACTTTTGC TAATAATAAA AAGATCACTA 105300 AATAAAAAG AAAATAACGA TAATGAAAAT GCTAATTTAG AAAATATACT AATAAGAAAA 105360 GATCTAAAAT ACTATGAAAA AATCATGGGA AAATCCCTAT TAATGCAAAA AATTTTTGAA 105420 CTTGTAATAA AAATAGCAAA ATCAAATGCA TCTATTCTTA TAACGGGCGA AAGCGGTGTT 105480 GGTAAAGAAA TAATAGCAGA TGCTATTTTT GATCTTTCAA ATAGAAATGA CAAACCATTT 105540 ATAAAAGTAA ATTGCGCAGC ACTTTCTGAA AGCATTCTTG AAAGTGAACT TTTTGGCCAT 105600 GAAAAAGGAG CATTCACTGG AGCAATTTCC AAAAAAAAAG GCAGATTTGA ACTTGCAAAC 105660 AAAGGCACAA TTTTTCTTGA TGAGATAGCA GAAATTTCAC CTGAAATTCA AGTCAAGCTT 105720 TTAAGAGTAC TGCAAAACAA AACTTTTGAA CGTGTTGGGG GAGAAGCTAC AATTAAAGTT 105780 GATATCAGGC TTCTGGCTGC AACAACAAA AACATTGAAG AGGAAATTAA AAAGGAAAAA 105840 TTTAGAGAAG ATTTATTTA TAGATTAAAT ATCATTAATA TAAACATACC GCCTTTAAGA 105900 GAAAGAAAG ATGATATATC TTATTTAACA AACATACTAA TAAAAGACGT CGCAAAGGAA 105960 AACAATAGAG AAGAAAAAAC TCTTTCTAAT GATGCAATGA AAGCTCTCTA TTATTACGAT 106020 TGGCCAGGAA ATATTAGAGA ATTAAAAAAT GTGCTTGAAA GTGCATTAAT ATTATCAAAA 106080 GGCAAACAA TCACTAAAGA AGATTTGCCA GCAAAAATCA AAAATAATGA AAATCTTATA 106140 TTTAAAATAA CACTACCAAT AGGAATTAGC CTAAAAGAAG CTGAAAAAGA AATAATAAAA 106200 CAAACACTTT TTCATTCCAA AAACAACAAA AGCAAATGCG CCGAAATACT AAAAATAGGA 106260 AGAAAAACTT TACACAATAA AATAATCGAA TATAATATTG ATTAATAGGA TTTATTTTAA 106320 ATTATTAAAT TATAATGGGT ACAAAAAAT AATACTGCTT TAAATTCCAT GTATATTTTT 106380 GAAACCAAAA AATTTTTTAA TGCCAATAAT TATATTAAAA TGAAACACTT TCTTTTAAAA 106440 TCATGGCGCA AAAGTGTAAA AATATTTTTA TCAAACAAAT AATTATACAC CATTATTTGT 106500 TAATAATCAA TACAATTTGA TAATTTAATA TATTTAGCTG GCTACAGAGC CTGACCTTAC 106560 TTTAAAAACT TTAAAGGGTT AATAGGAATA TTTTTTTTTA ATATTTCAAA GTGCAAATGA 106620 GGACCAGTTG CGCGACCCGT TTGCCCAACC nTTCCAAGAA ATTCTCCCGA TTTAACAAAA 106680 TCACCTATCT TTACAGAATA TAAATTTAAA TGCCCATAAA GAGATTTAAT ATTATTTTTG 106740 TGACCAACCA CAACAAAATT CCCATAAAGA TCATTGTATC CAGCTTCAAT AACTATTCCA 106800 GAAGAAGA ATACACTTCA GCATTCATTG GAGCTGCAAG ATCTATTCCT GTATGGAAAC 106860 TTTTGTTGCC AGTGAAAGGG TCATTTCTAA ATCCAAAATC AGAACTAACA ATAAATTTTT 106920 TTAAAGGAAA AATAAAATTG GCATTTAAGA AAAAAAGCAA TTCTGTGCCT GAAAAAAGTC 106980 CAAAATCTGG ATTCTTAACA AAATCAAAAA AATAAAATTC ATAAACTCTG TCGTTCCTTT 107040 TAATTTTTAC CTTTTCAGCT TTAGCAAGAT CCCTTGTTGC TAAAAGCAAA TTATTAAATC 107100 TATAATCTTT ACTATCAAAA ACAAAAACTC CTTTTTTACT GGGAATAAGA ATCTCTTGCC 107160

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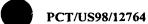
WO 98/58943

221 CCGATAAAAA ACTTCACTTA ACAGCTGGCG GGGCTGCAAA TATTGTCAAG CAAGCAAAAG 112500 TTTTACAAAC AGGACTTATC CATTTTAGTG AAAGATATAC ATTAAGAAAA GATCTTGAAA 112560 ACTTACTAAA GGAGGCAAAA TTGGAACATC CAGACGGAGA AATTTTTTTA ACAAGAGATG 112620 GAATGAGGCT TGAAGCAAAC AAAAATAACT TTATTATTAA ATAGGAGGGT ATATGATAAA 112680 TGTAGAAAAA GTTACTAAAA TGTATGGGCC ATTTACAGCA CTATTTAATG TTAGCTTTAA 112740 GGTTGAAGAA GGCGAAGTAC TTGGTATACT TGGCCCAAAC GGAGCCGGAA AGTCCACATT 112800 AATCAAAATC TTAACATCAT TTCATTATCC AAGCAAAGGT AATGTAAAAA TTTTTGGAAA 112860 AGACATTGTA GAGCATTCGA AAGAAATACT ACAGCAAATA GGATATGTTC CTGAAAAACT 112920 AGCTCTTTAT CCAGAGCTTT CTGTTAAAGA ATATTTAAAG TTTATATCAG AAATAAAAGG 112980 TGTTAAAAAA TTAAAAAAAG AAATTGACAG AGTAATAAGC ATATTCAAAT TAAAAGAGGT 113040 TGAAGATAAG CTGATTTCTC AACTTTCAAA AGGATTTAGA CAAAGAGTAG GAATAGCTGG 113100 CGCTTTAATA AACAATCCTA AACTTGTAAT ACTTGATGAG CCAACAAACG GTCTTGATCC 113160 AAATCAAATA ATTGAATTTA AAGAATTTTT AAGAGAACTT GCAAAAGAAA GTACAATATT 113220 ATTCTCTTCG CACATACTAA GCGAAGTAGA ATCTATTTGT AAAAGAATAA TTATTGTCAA 113280 CAACGGAGTA ATTGTTGCTG ATGACACAAA AGAAAATATT ATTAAAAATA AACTTAAAGA 113340 GATTGAAATA GAATTAATAG TTTCAAAAAA ATCTGAAAAT GAGAAAAAAA TTTTCAACAG 113400 CAAAAATGAT ATTTTTCAT TAATAAAGCT TGAAGAACAC GAAAAAGACT TAAATATTTC 113460 ATTAAAACTA TCTCAAGGCA AAACAGAAGA AGATCTCTTT AGCTACATAG TAAAAAATAA 113520 TATAATCTTA AAAGCAATGA TTCCAAAACA TGAAAGCCTT GAAAAGATAT TTAGCAAATT 113580 AACCAAGGAG AGAGAAAAAT GAAAATAGAT TTAAAGCAAT CTTTATCGCT TTCTAAAAAA 113640 GAACTAAAAA TATTATTTGG AACCCCAACT GCATACGTTG TGATGCTATT TTTTTTAATA 113700 TTCATAAACT TTTCATTAT TTTTTTATCA GGATTTTTTA TTAAAGACAA TGCATCTCTT 113760 ACCTCTTATT TCTCTTCAAT GCCTATTATT TTAATGTTGG TACTGCCAGC ACTTAGCATG 113820 GGAGTATTCT CAGAAGAACA CAAAACAGGA AGCATTGAAC TTCTTTATGC TCTACCGCTA 113880 AGTCCTCAAG AGATAGTCTT GGGCAAATTT ATTACGCTTA AAATATTTAC CTTAATACTA 113940 TTCTCACTTA CCCTACCTCT TACAATAATG ACAATTTTCA TGGGCGAATT TGATCTTGGG 114000 ATAATATTGC TTCAATATCT AGGAATAATT CTTTATTCTC TTTCTGTGCT AAGCATGGGA 114060 ACATTTATAT CCTCCATTAC AAAAAGCCAA ATAGTCTCTT ACATTCTTAC CGTATTTACA 114120 CTGATATTAA TACTATTTTC TGGGAAATTG GTTATGATCT TTGGAAAAGA AAATATAATA 114180 GGAGAAATAC TTAATTTTGT TTCAATAACC AATCACTTTA GCTATTTTAA TATGGGTATA 114240

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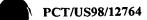
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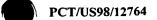
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GCACAAGCA	A TAGGAAAAA	TGGAGCCGTA	A TAAGAATTO	ATGCTTTATC	AATTGCAATC	143880
TATTCAAAA	C TTACAACAA?	AGAGCTAGGG	S ATGATGGATI	TCTCATATTC	CCCACCCTTC	143940
TCAAGAACT"	r gggatatati	· AAATATTGC1	GGCAATGCTC	CCAAATAGA	A AGAATTAAAT	144000
TAATTTAAT'	T CTTCATGCT	ATTGGTTGC	C CCGTACTTGA	A AAGAACATCT	CTCCAAAAAG	144060
AACCATTTG	G ATTAACCTT	A TTTCTGTCA	A TTACTGCCAT	CTTAATAGG	r ATATGAACAA	144120
ATTTTGTAC	T CCATAAACT	A ATCAACATT	r TTGTCTTACO	AGCCATTGC	A GCATGCACAG	144180
CATTCGACC	C AAGCCTAGC	A CAATAAAGC	AATCACTGG	C ATTAGCAGG	r GAACTTCTAA	144240
TAATATAGC	T GGGATCAAT	G TATTTAAGA	G TAAATTGTA	r ATTTTTTGC	TTAAAATTT	144300

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		7		239			
C'	TGTAATTTT	ATCTTTAATA	TAAAGCCCAA	татсстсата	AAGCAAATTC	CCAGAATCGT	144360
C'	TTTCTTCTT	AGGAAAATGA	TCAAAATATT	TTTGGCCTGC	TCCTTCTGCT	ATCAATATTA	144420
C	TGCATGGGG	AATCTCTTCT	AAGCTTTCTT	TCTCTAAAAG	TCGTCTTTCA	AGATGAACAA	144480
G	AAATCCATT	AGGACCTTCT	ATGTCAAAAT	CAAGTTCTGG	GATTAAACAA	AAATTAACAT	144540
С	ATTAGAAGA	AAGTGCGGTA	TGAGCAGCAA	TAAAGCCAGA	ATCCCGTCCC	ATAACTTTAA	144600
С	AAGTCCAAT	GCCATTATAA	GCACTATTAG	CTTCAAAATG	AGCACCAGCA	ACAGCTGCAA	144660
C	AGCTTGTTC	TACAGCAGTC	TCAAATCCAA	AAGATTTTTG	AACAAACATA	AAATCATTGT	144720
C	TACGGTTTT	AGGAATGCCC	ACAACTGCTA	TTTTTAAATT	TCTTTTTCT	ATCTCCTCAG	144780
C	AATAAGAAG	AGACCCCTTT	TGAGTACCAT	CCCCGCCAAT	GTTAAAAATC	ATATTAATGT	144840
T	CATTCTCTC	TAAAGTATCA	ACTATTTCCA	CAGGCTTAAT	ACCACCCCTT	GAAGAACCAA	144900
c	SAATAGTACC	тссаалтта	TTAATATCAT	CAACAACATC	TGGATTAAGA	ттаатаааа	144960
C	GTGAATTTGA	CTCAGGAAGA	AGCCCTTGAT	ATCCAAATTT	TACTCCATAA	ATATTGCGAA	145020
C	CCCCATATAT	TTTCCATAAA	GTTCGCACAA	TAGAGCGAAT	· AACATCGTTA	AAACCAGGAC	145080
1	AAAGCCCACC	CACAAGTAGTA	ATAGCAGCTI	TAACATGCCI	GGGCACAAAA	TAAATTTTTT	145140
(CTCTAGGCCC	C AGCTTTTTCT	AAAAGAACAT	CTTCATACCI	ATCTCCCTTA	A TCCTCATTCC	145200
,	гататасас	r aaacttgatt	TTATTTTTT	CATTAACAAA	A ATGGGAAGA	A CCCTCACTAG	145260
(CATAAAAAT(C AATCAAAGGA	A TTGTTTTGCT	TGCATTCTCC	CAAGCTATC	TAAAATTTTA 1	145320
,	СТАААТТТТ	C ATTTTTAAT	CTATACACC	A AATACTCCTT	r TATAGAATT	A TAACCTAATT	145380
	АТТТТСТАА'	T AAATCGACT	T TGATCTTTA	A TCATATCGT	A TATGTCATC	G TAAATATAAG	145440
	GAGACCCTT	C AATAGGAGA	AAATTAATT 1	TACCAGCTA	r GAATTCAAA	А ТАТТТАТТСА	145500
	ACTTTGAAT	т тттстсааа	A TCAATAAAT	G GAACTCTAT	r ATTAATAGC	C TCTCTGAAAC	145560
	TTTTTGCAA	A AGGCACAAA	A CCTATAAAC	r ctattggta	T ATTAATATT	A TTCTTAACAA	145620
	САТТААТСА	A ATTTTCACA	C ATAGCAATC	T CTTCACTAG	T TTCTATTCT	A TTTAGCACCA	145680
	CTCTAGGAT	ТТАТТАААА А	C ATCATCCTC	т таастттса	A AGAGGAACT	C AAAGAAATAA	145740
	GTTCAATCC	C AACAACCAA	а тстттааат	C CAAGGTTTG	T CCCCTCAAT	C TTATCTTTAA	145800
	AAAAATTAC	C AATATAATC	C CGTTCGGGG	C TTTTTTGCG	G AAATCCTAA	A TATAAAAGAC	145860
	GATAAAGAG	GC ATTCTTAA	A AAAGAATAA	.G САТТААСТА	T GGAAGGGGT	T TCTGGTATTG	145920
	TAACAATTA	AC ACCGCTGTA	A GATGCCAAA	т ааааатста	TGTATTATA	A GAAGTTCCAG	145980
	ATCCCAAAT	гу тааааат үт	'A AAATCAGCA	A TAAGATCTI	T TTGAATGG	т тстатаатст	146040
	TTTTCTTA	AT AGAAAAAGG	GA AGATTAGCI	G TTCCCGTAT	AAGAGCATO	CA CCTGGAATAA	146100



GATAAAGCTT ATCATAAGAT GTTTTACATA CTAAATCTGA AAAACTTTTA CTCTTTTAT	146160
TAATAAAAGA ACCAATGCCC ACACCCTTAT TTTTAACCCC CAAACACGTA TGTAGATTAG	146220
AGCCACCAAG ATCAAGGTCA ACAAGTATTA CAGTTTTACC CAAACTAGAA AGCTTATAAC	146280
CAACATTTGC AACAAAAGAT GTTTTTCCAA CACCGCCTTT GCCACTTGCC ACAGGAATAA	146340
TTTTAGTCAT TCTTAAATCC TAATTATCCT TACGATCTTT TTGAAAAATT TTCATAAAAT	146400
TGAAAATCCC TAAAAATCTA GATTTTTTCT CAGCATCTTT ATTTAAATTT TCATCCTCTT	146460
TAGAGCCTGA AATTAAATCT TTAATTAAAT CTTTATTATT TGTAAAATTT TCAATGCTAT	146520
CTGGTTTACC ACAAATTACA ATTTTATCAT CTTTTAAAAA AAAATAATCG CCATCAACAA	146580
ATTCATACCT AGAATTACTT AAATTTCTAA CAGCAATAAC TGTAATCCCA CATTCTCTTC	146640
TAAGATCGGC TTCAAAAAGA GTTTTACCAA CATATTCTTT GGGAATAACA GTTTCAGCAA	146700
CAATAATATC ATACCCAATA ATATTATAAG TTGAAAGATT TGGAGATACT AATAATGGAG	146760
TTAATCTTCT TGCAGCATCT TTACTTGGAA ATATAATTTT TGTTGCCCCA AGAGTTTTTA	146820
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TACAATAGTG AGTAACAAGA GCACTTTTGC CAAGATCATC ATCAAAATCA ATAACAACAG	146940
CGTCTGTATC TACTGGAATT ATTCTTTTCA AAGCATTTTT AGTGAATTGC TCAACAACAA	147000
AGCTTTCTGT AGATATCACA TCATATTCTT CAATAAGCTC TTTAGATGTA TCTATAATAA	147060
TAATTTGACA ATCAAGCCTG CTTAAATCTT CAAGTAAGTG AATGCCTAAA TTACTAAGTC	147120
CAATAATAAC AAATGTTTTC ATATGCTTCA ACCAACCAAA ATATCTTGCC TTGGCCTTGT	147180
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TCCTGCAAAC ATAGTAAAAA TTATAATGAC TTTCCCCCAA AATGACAAAT CCTGAGTTAC	147300
TCCAACTGAA AGACCAACCG TTCCAAAAGC AGAAAATACT TCATAACCTA AATCAATAAC	147360
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TGGTAGAGAA ATTATTTGAG TTCTTCCGCT TATTAAAGAA TTATCAAGAT AATTAAAACC	147660
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PCT/US98/12764 WO 98/58943 241 AGTGTTATTT ACATCTCTAT AGACCATAAA CCCAAGCCCA CCACAAATTA TTAAAATAGA 147900 147960 GACCACAACT ATAGCTTCAG GAACATCTCG CCATGCATAA ATACTCTCAG AATGCATGGA AAAACCTGCA TTGCAAAAAG CAGAAATTGT CGTAAACAAA GCCTCTAAGA ATGAAATATT 148020 CACTCCCCTA AGTTTAAAAC AAATAAGTAT TAATATTAAA CCTATCATTT CAATTGAAAA 148080 AGTTATAAAC AATATGCTTT TTAAAATTCT AATAGGATTA TATTCTATAT TTGAAAGGGA 148140 ATACTGCTTT ATTATTCTTG CATCTGTTAA ATTCATTTTC TTTTTAGGTA TAAGCAAATA 148200 AAAAGTAGTA ATACTTATAA ATCCAAGTCC CCCAAGCTGG ATTAGCAACA TTATCAAAAT 148260 AAATCCAAAA GTAGAAAAGC CTTCCATTTT AACCGTTGTA AGGCCCGTAA TACTTACAGC 148320 AGAAACAGCA GTAAAAAGAG CATCAATGTA TGCTAATTTG CCATCACCTT CCCAGGAAAT 148380 AGGCAACATC AACAAAAGAG AGCCTATAAA CATAATTAAA ACAAAATAAC TAAAAAGTAA 148440 AAACCTGTCG CTAAATTCAA ATTTCAACAT ATCATACAAA AAGTTGTTTA AATTATTAAA 148500 AATTTATCTT ATATAGCATA ATATTTTAAC ATTGAAATAT TATCATAATT ACATTATTTT 148560 TAATATGT TTGAAATAGA ATCAAAAGCA TTTATTCCTA CAAAAGAGTT AAAAAGAATT 148620 ATCAAGCTAG CAAATAAAAA ATTTAAGTTT ATTAAAGAAG AAATAAAAAC TGACATYTAT 148680 TACTCAAACC AAAAAAAAT TATAAGAATA AGAAAATTAA ATACTCTAGA AAAAATTGTC 148740 ACATTCAAAA AAAAAATATT AGACAACAAC AATACTGTAG AAATTAATAA AGAGATAGAA 148800 TTCAAAATAG ATAGTATTAA TAATTTTTTA ACCCTTATAA AAGAGCTTAA ATTTAAAAAG 148860 CTATACAAAA AGATAAAAAA AAGTTTAATT TATCAAACTA ACAATTTAAA TGTAGAGATA 148920 AACGAAATAA AAAATCTTGG GTTTTTTTTA GAAATAGAAA AAATAATTAA CAATCAAAAT 148980 149040 GATATAGACT TGGCAAAAAA AGAAATTGAC AACATAATCA ACCAATTTGG ATTAAAAGAA AACATTGAAA CTAGACCTTA CTCTGAATTA CTTTCATTGG CAAATCAAAG TAAAAAATAA 149100 TTCATTGGAA TTAGAGCTTA AAGTAGAGAT TACAAGCCCT TGATTGCCAT AAATTCCAAT 149160 CTGAGGGCTT TTAACATTAC TCTTAAAATT CTCAAGCTTA TTTAAAAAAT ACCAATTTTT 149220 ATTCTTAAAA TAAATTAATC TCACATTATT ATTGTCCTCA AAAGCTAAAA ACAAATTATT 149280 TTTATAAAGC CCAATGTCAG CACTTAAACC TTCCATTTCA ACATTAGGAC TTATATAAT 149340 CCATCTACTA CTTTTCAAAG GACAAATGTT TACAATAGGT CTATTTTCAG AAACAAAACT 149400

CATAATTATT TGATTAAAAT TAGAATCAAA AAAGCCTTTA ATAAAATTGG CCATATAAAC

AGAAGGAATA TTTGCATTTA CCCAAGCATT TTCATTGTTT ACAATAAATT CAGATTTAAT

CTCATTATTT GACTTATAAT TATAAAAAAT GCCCAAAAAA GGTTCAGATA TTAAACCAAT

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149460

149520

149580



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AAGCTCGCTG	GCAAAATCAG	CCCCTGATTT	CGTAACAAAA	GCAATATATA	AATTTCCTTT	149880
AGAATTAATT	GAAAAATCAA	AATTAACTAT	ATTAGTAATA	TTTCTATTAA	CAGATGAATC	149940
AAGATTAAAC	CAACCAACAT	CCTCAATAAA	TTCAGCAACT	TTAATATCAT	CGCTATTTTC	150000
TAGCTGATAA	GCAATATAAA	TATTGCTTTT	ATAAATCCTT	AATACATATT	TTTTAAGCTT	150060
GGCAGTTAAA	TTTAAAACAG	GCAAATCTTT	TAAAGTAAAA	AATAAATCTT	CTTTTTCAAT	150120
CTTAGATGTT	TTAGCTTTCA	GGGAATCGCT	ACTTAAAATT	GAAAAATCTA	AATCTGTAAG	150180
CGAAAACTTT	ATGTTTTCAT	TTTTAGTACC	AACATATATT	ATTGCATAAA	GAGAATTTTT	150240
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ATCGGAAGAT	TTGCTAACTT	TTAAGTAAAC	ACTTCCTTTC	CCATTTTTGC	TTAAAATACT	150420
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GAACAATAAA	TAAAAATTTA	AATTTAAAAG	TTAAAATGTA	ACTAATAAAA	TTTTGAACAG	150660
ACTCTGTAAA	GTTGAGTTGA	TTTAAAGTAT	AAAATAAAAG	GCTTGCAAAA	GTGCAAACAG	150720
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TAATTAAATT	TCTAACAAAC	CCAATAGAAA	TTTCACGATA	AATAAATAT	АСАААААААТ	150840
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CCGCATAAGG	ATCTAAAATT	TTACCTACAT	TGCTAACAAG	ACCATATTT	CTTGCAAGAT	150960
AACCATCAAT	AAAATCAGTA	AATTCATTAA	. ААТТААТТАА	AAACCAAATA	ATTCCAAAAA	151020
ACAAATACGA	ААААААТАСА	ТТТТССАААА	TAAAATAAAA	TAATATGATA	AAGGAAAGTG	151080
CAATTCTAAC	TAATGTTATT	TTATTAGGGG	TAATGACCTT	GATTAAATTA	TTCAATTTAT	151140
CAAATCTCCT	ТАТСТСТТАТ	' TTTAAATAA	AATTTAAATTA	GAGCTTCATC	AAGTTTCATT	151200
ССАТТТАТТТ	GCTCATTTGT	TCTTGTTCTA	ATAGATATTC	TCTCTTCTGT	TGCTTCTCTC	151260
TCACCAATTA	ТАААСАТАТА	AGGTATTTT	TTAGCCTGAT	ATTCTCTAAT	TTTAGCATTC	151320
ATTCTTGAGG	AACTATTATC	AAGCTTTATT	CTAATCCCCT	CATTTTAAA	A TTTATTAAAA	151380

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						-
GATAACCATA	AAGGAAAAGC	ACCACCATAG	TGCTCTACAA	GAATTCCAAA	AAATCTTTCA	151500
ATAGATCCCA	ACAAAGCTCT	ATGAATCATA	AATGGTCTTT	TTTCTTTACC	ATCCTCAGCG	151560
GTATAAGTCA	TATTAAATCT	CTCAGGGAGA	ТТААААТСАА	ATTGAATTGT	ACTCATCTGC	151620
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CCACCCTTAT	CAATTTCATA	AGGAACTTCA	AAATCGCTTA	AAGTCTCTTC	AAGAACTTTT	151740
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GCCTTTGGGT	TGCTAAAGCC	AAATTTACTC	CACATATAAA	TAGCAAACCT	AAGAACTTCT	151860
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CCTCTGGCTC	TCATCATACC	ATGCAAAGCA	CCTATCTTTT	CATAACGATA	CACAGTGCCA	151980
AGTTCGGCCC	ATCTAAATGG	CAAATCTCTA	TAAGAATGCT	TACCTGTATT	GTAAATTGCA	152040
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CCAATATGAG	GAGTAAAAAG	AATATCATAC	CCATTTTTGG	AGTGCTCTTC	TCTCCAAAAA	152220
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ATTCCAACAT	TAAAAGAAGT	ATGAAAAAAT	AAAAGTCCCA	AAATTCCAGA	TATTACTAAG	153240
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AATAATATTA	AAATAGTGCT	AACACCCAAA	AACCCAAACT	CTTCGGCAAG	AATAGAAAAA	153360
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CCTTTGCCCA	AAAGACCGCC	AGAACCAATT	GCTATTTTAA	CCTGATTTAA	ATTCCAACCA	153480
GCACCCTTAG	CATCAATAGC	CGGATCTAAG	ААТАССАААА	ACCGTTTAAT	CTGATAAGTC	153540
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АААААТАСАТ	ААААТАААТ	ТАТТТТААТА	CTCAAACCAT	ATTTAGAAAT	GAAAAATCCT	153660
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AGTAAGCTTA	TCAAAGCCAA	ATAATCATAA	TTTTTTCTAA	AAACCATTAA	TCTACCTAAT	154380
ATACCACGGC	CTATAACCTT	TAAGAATATC	TTCATAACTT	TGATTTGCAA	AAATGCCTTG	154440
САТТАТТААА	TCTGTAGATT	TTGCAGGCCA	CCAATCCACA	TTACTTTTTG	CCTCAACCAA	154500
ACTAAAAACA	ATAATTTGAT	TATCAGCTGA	ACCGTTATAA	GGGGCAAGTC	CAATAAAAGA	154560
ACTATTTTCA	AAACCATCTA	TTCCAGTTTG	ACCAGTACCT	GTTTTTCCTC	CAACCTCAAC	154620
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TTTCAGAAGT	TTAAATGTGT	ТТТТАСТААТ	AAGATTTGTC	TTTCTTAATA	TTTCTGGTTT	154740
ATTTTCAAGA	ACAACCTTAT	TAGTACCACC	ТТТТААААТТ	TTATTTACAA	TTCTAGGTTT	154800
ATATACAACA	CCTTCATTTG	CAATCATAGC	AACCATATTA	ACAATCTGCA	TAGGAGTAGC	154860
АТТТАААААТ	CCTTGACCTA	TTGAAAAATT	TACAGTATCT	CCTCCTACCC	AAGGCTGATT	154920



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	7		245			
AAAAGTTTTT	TCTTTCCACT	CAGGACTAGG	AAGAAGGCCA	GCTACTTCAT	TTGGCAAATC	154980
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AAGATACTTA	AGCCCAAGTG	ТАТАААААТА	AACATTAGAA	GAATGTGCAA	TCGCCTCTTC	155100
ТАААТТААСА	TACCCATGAC	CTCCGGGCTT	CCAGCAATGA	AAAATTCTAT	TTCCAACTTT	155160
AAAATATCCA	GGACAATAAA	TTTTACGATC	TTTGTCTATA	ACTCTTTCTT	CAAGAATGGC	155220
AGCAGCAACA	ACTAATTTAA	AAATAGACGC	AGGCGGGTAA	ACAGATTGAA	TTGCTTTATT	155280
TAAAAAAGAG	TAATCTTCCT	TATTATCTTT	ATTGTAAACA	TCTTTCATAG	AATAATAAGG	155340
ATAATTGTGA	AGAGCAAGAA	CAGCACCTGT	TGATGGTTTT	AATACTACAA	CAGAACCATA	155400
CCTTTTGCCT	AAAGCATTCT	TAGCAAGATC	TTGAATATCT	TTATTGATAT	TAAGCACAAC	155460
ATCATTACCG	GGCACCATAT	TTTTTATAAT	AGAACCATCG	TCTATTCTTC	TCTCCTTAGA	155520
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AACGCCCAAC	TTTCCAATCG	TAGAAGTATT	ATCATACCCA	CTAACATTGT	AAAACGTCCT	155640
AAGTTCTCTT	TGATTTATTT	GCCCAACATA	ACCGATTGAA	TGAGAATATG	AATCGTCAAC	155700
TAAATAGTTA	CGCTTAAAAG	AATAGGTCCA	CAAAAGAGCA	GGATAATAAA	ACTTTTTTC	155760
AGAAATTTTG	AAAAGCATCT	TTGGGGTAAG	ТТСААТТАТТ	TCAACATCTT	TAAGATATCC	155820
ACCAGGCTCT	TGAAGTTTAG	ACAAAATAAT	TGATTTATCA	ATATCTAGAG	TGCTTGATAA	155880
AAAATCTATC	ATCTCAATTC	TAGTAGCAGC	AGGCATATTG	TAATACTGTT	GTAAGCTTAT	155940
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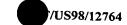


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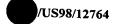
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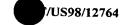
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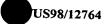
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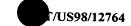
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TCCCTTTTCC	TCTAAAATAG	TATACTCCCC	CATTAAAAGT	AAATTTCCGG	GTACAGAAAA	182040
ACTAATCAAA	TCCATTCTAA	GTCACACCCA	ACCTTTGAAA	CAATAAAATC	AATGCCAGTA	182100
AAATTTTGCT	TAAGTCCTTT	TAAAATAGTA	TTTAAATTTT	CCTCCAAACA	AAGAAACTTT	182160
ACTTGGGGGC	CTGCATCCAT	CGTCTCAAAT	ACAAAAATCC	CCTCATTTCT	CAAATCAGCA	182220
GCATACCTAA	TTAAATCTAT	TGTACTATTT	ТТААААТААА	AAATAGAAGA	TGCAAACATT	182280
AAGGCAAACA	TATTCTGATA	ACTTTTTACA	ATAGTTGCTC	CAAAATGTAT	AAAATCCTTT	182340
TTTTAAAAAA	AAATATAAAG	CGTCTTTAAA	AATCTTTTTA	CTAGAGGCAA	TCCAAGCATC	182400
АТААТАААТ	TTATGCCGTT	TGCAAATATT	CATTGCGGCT	CTTGAAGACA	ATTCTTTTTC	182460
ATTACTATCA	ATTATGGCAA	ATATTATTCG	CAAATCATTA	AAATAAGATT	GATCTCTTAA	182520
TTGAAAAGAT	TCTTTTGAAC	CCTCTTTTAA	AATAGTAAAC	CCCCCGTAAA	TAGCCCTTGC	182580
CGCAGAAGCC	GATCCTACTC	TTGCAAGATT	AGATGCGCTA	TTACAAGAAT	ATTTATTAAA	182640
ATATTTCAAA	ATACAAGCAG	CAATAGAAGC	AAAACCTGAA	CTTGAACTTG	CAAGGCCTGC	182700
TGCTGTCGGG	AAATTGTTTT	TACTTTTAAT	ТТТАААТСТА	ACATTTGGTT	CATTAAGAAT	182760
TTTTCTTGCA	TAATCAAAAA	ACACCTTTTC	TCTATTTTTT	AATATAACTG	GCTTTGAATT	182820
ТААААТТАТТ	TCATCTCGAT	TTGAAAGTTC	AAGCTCACTT	ATTGAATAAA	ACTTGTCAAC	182880
ACTAACAGCA	AGACTGGAAG	TAGCTGGAAT	GTTTAAAAAA	ACATCCTTTT	TCCCCCAATA	182940
TTTAATTAAA	GCTAAGCTTG	CATGAACTTT	ACACTTTATT	TTCATTCTCT	AACCTTATTT	183000
TCTTCAAAAT	TTTAAAAGCA	AAATCAAAAG	AATAAATATT	CATTCTTTCC	АТТТССААТА	183060
ATAATTTGTC	TTTTTCAAAA	TCAGAAATAT	TATATTTTGT	CTTTAAAAGA	TGGAGTATTT	183120
TATTAACATG	CAATCTCATA	TGACCCTTTT	GAATCCCATT	AAATGCAAGA	GCCCTTAATG	183180
CTGCAAAATT	ACTAGCAAGT	CCAACACAAG	AGAGAATACC	AATAAATTCG	CTCTTACTAT	183240



			261			
TTACATTCAT	AAATTTTAAA	CTTAAAATTG	AAGCTTCATT	ААААСАТАТА	ACCCCACCTT	183300
TAGTTCCAAC	TTGCAAAGGA	ATTTCAATTT	' СТССААСТАА	AGCATTGTCA	GTAGTATAAA	183360
ATTTACTAAC	GGGAAAATA1	TTGCCGCTTT	TTGAAGCAAA	TTTATGAACA	GAGGCCTCAA	183420
GAGCTCTTGT	GTCATTAAAA	GTTGCAAGAC	ACACCCCTGT	AATTCCATTC	ATAATACCTT	183480
TATTATTAGT	AACAGCTCGC	TCCTCTTCGT	AAAAACCTAT	ACTAGAAATA	AGTTCAATTT	183540
TTTTAGCCAA	ATTCCAAGAA	TCCTCTTTAC	CCGGTAGCAA	ATGCTTAAAA	TCTAAAACAA	183600
AACGGGCTTI	GGCTGTAAAT	TCACTAATAT	CATTGCTTAA	AACTTTTAAA	ACACACTCAT	183660
ATCCGAATTC	ТААААААТА	AATTCTGCTA	CACGCTCTGC	AATTGAGTTT	AGCAAATTAG	183720
CACCCATAGO	ATCACAAGTA	TCCACATAAA	TATTTAATTT	TTGAATACCT	AATTCTTTAA	183780
TGTGCCTAGT	TGACAACCTT	CTAAATCCAC	CCCCCTTTG	ATTCATATTG	GTTAAAAGAG	183840
GTTCAATCCA	AGTCTTAATT	TTATCACCAA	GGTCAACAAA	AATTTTACTT	AAATCTTTTT	183900
CCGATTTTAT	ATAAATTTGC	GAAATTCCCA	ACACTTCACC	CAAAGAATAC	CTTAAATCAG	183960
CATTTTCAAG	AATCTTTGCC	GCAAAATTTA	GGGCAGCAAC	AACAGAAGAT	TCTTCTGTTG	184020
CAATTGGCAA	AGAATAGTAT	TTGCCATTTA	ТТТТСАААТТ	TTTTACAATT	CCAATAGGAA	184080
AAGATAAATA	TCCAATATAA	TTTTCTATCA	TATTAAAAAG	AAAATCTTCA	TTGGCATTAT	184140
ТАТАААААА	ATCTTTATAA	GATAATTCCA	AAAAACTTTT	TATCTCTTGC	СТТТТТТСТА	184200
AAACGCTTTT	ATGTCTAAAA	ТТТТТАСТАА	GTTCCATAAA	ACTGCTTAAA	GACTCCAAGT	184260
TCATCAAGCA	AATAACTACT	TAAAAAATAC	TTATTATTTC	TAAACTCTAA	TAAACTTTTG	184320
CTTCCACTTA	AAAACATAGA	CATTTTTAAA	ATATGTTCAT	AATCAGAAAA	AAGACCAAAT	184380
ACAGCATCTT	CTCCTGAATC	ATAAAAAGCC	CTAAGAACAA	CTGCTGCAAC	ACCTATAAGC	184440
CTGGCTCCAA	GGGCAATGCC	TTTAGCAATA	TCCATGCCCG	TCTCATATCC	ACCAGATGCA	184500
AAAATATTAG	CCTTTAGAGA	ATCATCAATA	CTAAGTAAAG	TAAAAACCGA	AGGTATACCC	184560
CAATCAGAAA	AACAAGATGC	AATGTTTAGA	ТТАТТАСТСТ	TCATGCCTTC	ТАСТААААТС	184620
CAATTAGTTC	CACCACTCCC	TGCAAGATCA	ACATAAGAAG	CACCAAGGCT	GAACAATTCC	184680
TTAACGTCTT	TTGGCGAAAT	TCCAAAACCT	GTCTCTTTAA	CAATCAATGG	AACACTTAAA	184740
AAGTCTGACA	ATTTGGCTAT	TGACTCTCTT	ATTCCTTTAA	AATTTCTATC	TCCATCAACC	184800
TTCATCAATT	CTTGTCCTGC	ATTAAGATGA	ACAATAATTG	CATCAACTTC	ТААТСТТТТА	184860
ATCATTTCAG	CTATTTTAGA	AATACCAAAT	TCAACAATCT	GAACAGCACC	AACATTGGCA	184920
AACAAAGGAA	TATTATGAGC	ATACCTTTTA	AGAGTAAAGT	CTCTTATGTA	CTCGGGATAC	184980
ТТАААСАААА	GCTTAAAAGA	ACCTAGCCCT	ATAGGAATTT	TTAAATAATT	TGCAATTCTA	185040



ACTAAAGA'	TT TATTAAAGTO	C ATTCCCCTC	TTACTGCCC	CTGTCATGG	AGAAATAAAA	185100
ACAGGCATA	AC TAATATTGT	A TCCAAATATO	C TCTTCTTTT	A TGTTTATCTC	GGAAAAATTA	185160
AAATCACT	AA GAGCATTGTO	G TTTTAGCTT	A ATAAACTTT#	A AGAAATTACA	GCCACCTTTA	185220
ACATCGTT	TATTTAAAC	A AATCTCAAT	A TGCCTTTTTT	TATTTTCTAA	TATATTAGGC	185280
TCGATACCO	CA TAAACTCGGT	T ATCCATCATT	CCTTAGTTCT	TTTAAATAA	ATCCTCTGGA	185340
TTCACCAGO	GA ATTATTTAT	TTTGAAAAA	ATCTTTATAT	TCTTCAAAAT	TTGCATTATT	185400
TCTATTTT	T ATAAGACCTI	CAAGATCCCA	ATAATTTAAT	ACATCAAAAG	CACTCTTTTC	185460
GATGGTAAC	ЭТ ТСАТАА <u>А</u> ТАА	TCATAATATI	GCCAGATCCA	TAAGAGCAAA	ACAATATCTT	185520
TTCCCCTGT	АТАТСТТТСТ	TGGAAAATAC	TCTTTTTAAA	TAAAATGCTA	AAGATAGAAA	185580
AATTGAACC	т статасааат	TTCCCACTTC	CATAGCAGCT	TCAACTCCAT	CGTAAAAATC	185640
TATTGATTC	т ааатаассат	' TTCTAACAGA	TTCGTCATCG	СТАТААТАТТ	ТТТТСААААТ	185700
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GTTTGCATT	A TTGTAACATT	CAACTGAATA	CTGACCTCGC	ACCTTAGCCT	CAACACTTCC	185880
AAAAGGCCT	A AAAAAATCGT	CAACATCATC	AGTATAAACT	CCAAATTCAG	ATAAATTGAT	185940
CGAAAGTAG	C TTTGGATTTT	TTTCAATCAA	AATTGCAGTT	GCGCCGGCTC	CTTGGGTAAT	186000
CTCAGCCGT	A GTAAGATTGC	TATAATGTGC	AATATCTGAA	GAAAAAACTA	TGCCGTATTC	186060
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ACACGCATG	T TGAACCTGGA	AAGTTAGAAA	ATTATTTCCC	AGACAAATAC	CAGATTGCTT	186180
TAAGGCTCC.	A AAAACATAAG	AAGAAATGGC	CTTTGAATGA	TCAACGCCTG	TTTCAGTTCC	186240
ACCCAAAAG	Г АТТСТААТТТ	TGCTTAAATC	AAGATTATTG	TTGTCAAAAA	TAAGCTTAAC	186300
AGCCGAACT	r GCCATGGTTA	CACTATCCTC	ATTAGGACTG	GTAAACCTAA	AACCTTTTTG	186360
CAAGGTTGC	A TCTATTGCTC	TATTGATTTT	TTTAAAAAAA	ACTTCATTAG	AAAAAAAA	186420
AGGATTTTC	C AAAAGAACAG	АААААТСТАА	ATAATTTAAA	GGTAAAAAAA	TTCTAATATC	186480
ACTAATACC	T ATTCTCATAT	ACTCCTCAAT	GAATTAATGG	CCTTAAGTAT	TATATTATAA	186540
TTTACAAAA	A TTAGCAAAAT	СТТАТАТААТ	ААААССТААА	AATGGAAGTT	TATGAAAATA	186600
GCCGTGCTT	TATCTGGAGG	AGTCGACAGT	TCTGTTGCCC	TTTATAGAAT	TATAAACAAA	186660
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ATTGGAAACT	GCCCTTGGCA	AGAAGATTTA	AATTATGTTG	AAGCTATATG	CAACAAATTT	186780



			263			
AATGTACCGT	T ATGAAATAA1	· AAACTTTCAA	AAAGAATATT	ATAACAAAGT	AGTAAGCTAT	186840
				ATATTTTTTG		186900
ATAAAGTTTC	GAGCATTTT	TGAGAAAATC	AATAGCCAAT	ATGATTTGGT	TGTAACGGGA	186960
CATTACGCTA	АААТАСАААТ	' AAAAGAAAGT	AAATTTTTAT	TAAAACAGGC	AAAAGATAAA	187020
ATTAAAGACC	AAAGCTACTT	TTTATCTCAT	CTCTCTCAAA	AACAAATGTC	AAAACTATAC	187080
TTTCCCTTAG	GCACATTACT	' TAAAAGCGAA	GTAAGACAAA	TAGCTAAAAA	САТАААТТТА	187140
CCCAACAAAG	ATAGAAAAGA	TAGTCAGGGT	ATTTGCTTTT	TAGGAAAAAT	ТАААТАТААС	187200
GAATTTATCA	AATACCATCT	TGGAGAGAAA	AAGGGAAATA	TAATTGAAAA	AGAAACGGGA	187260
AAAATAATAG	GAATTCACAA	CGGATATTGG	TTTTTTACAG	TTGGACAAAG	AAGAGGAATA	187320
AAACTTAGCA	ACGGGCCATG	GTTCGTCATA	GAAAAAGATC	TGGAAAAAA	TATTATATAC	187380
ATATCCCATA	ACGAGAATTA	TTTAAAACAA	GCAAAACGCA	AATTTTTAGT	TCACGAAATA	187440
CATTGGATAA	ACGACACACC	TACGAACTTT	GAAAATTTCA	AAATTAAAT	AAGACATGGC	187500
GAAAAGAAAT	ACTCATGCAA	АТТААААСТТ	ATTACAAATA	ACTTAATGGA	AATTTCTTTA	187560
AACAAAAAG	ATCAAGGAAT	CTCCCCAGGA	CAATTTGCAA	TTTTTTATAA	AAACACAGAA	187620
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ACCAGAAGCA	AGTCTCTCAG	AATATTCCAG	TATCTTATTC	AAAGAAGAGC '	ГТААСТТТТТ	188580



CACAAGATAA	AGAGTTGCAA	TAGCAAGCAT	AAGTAATGTA	AATACAAAAC	TAATTGCTAA	188640
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GCTAGAGAGT	ATTCCAAAAT	TAATCCCTCT	' AAAAGACCTA	ТАТАТАТАТ	CCATTGAATA	189000
AAGAAACATA	AAATAACCAA	AAATGTTATC	TGTTTCAAAA	TCTCTTAATG	GCATACCTAT	189060
TAATATGTAA	GGAATCTGAC	CTTTTTTATT	TTTTATCTTG	GAAATATCTT	TAAGTAAAGA	189120
CTCCTCAATA	GACCCGGGAT	CTGCCAACAT	GACAAAGGAA	TTGTAAACAA	TACTATTAGA	189180
CTCCTTAAGT	TTTGTAAAAT	ATTCTCTATC	CCCAATAGAT	TTGCCAAAAT	CACTATTATC	189240
CTTAACCGCC	GTAGTAAACA	СТАТТТТАСС	TTCTTTGTCT	GTAACCATCA	TATTACTACC	189300
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ACTAGCCTCA	TTTAAATTT	CAGCAGAGTT	GAAATACATG	GAACTAACCC	TAACTTTCTC	189420
TTCCATTGAA	GAGATGTAAA	GATTAAAAGA	ACTTTTAATG	CTTTCAATAA	GATTTATCAT	189480
AAGATTAAAC	TGTTGATCCA	CCAATTTACT	ATTAATAAGC	ATTCCAAAAG	CAAAAAACAA	189540
AATTGATATA	AAGAATGCTA	TCAGAATAAG	AACAAGTAGC	AACATCCTAG	CTTTAAGCTT	189600
CATACTAACC	ACCTCTTTTA	САААААТАА	ATTCTAAAAC	TCTGAAAAAT	CATCATCAAA	189660
ATTTAAATCC	TTATCAGCAA	TATCGATAGC	TTTTTTAGGA	TCAACTCGCT	TATTAATAGT	189720
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AGTAGAAATT	CCATTGCTTT	TCAAATTTTG	ATTTTCATCT	TTAAAAGAAT	TTTCAGGACA	189840
ATCTATTAAC	CTGAAATCAT	AATCATCATT	TTCTGGATTT	TCAATTTTAG	ААТСТТТААТ	189900
TTTGAAAAAT	AATACAGATT	TTCTAAGTTC	CTTAGACTTT	TCTAACATTT	TATCGGACAT	189960
ACTAGAAAGC	TGCTCACTGC	TTGAAGCTGA	AGATTGAACA	ACTTCTCCAA	CCTGATCTAA	190020
AGCCATTTTA	AATTGAGCAA	TCTGATCGCT	TTGCTTAGAG	CTACCTTCTG	AAATCTTCTT	190080
AACAAGATTA	GCCGTTTCTT	CAATTTCGGG	TAGCATTTCT	TTAAAGATCA	CTCCCGCTTC	190140
AGTTGCTACC '	TTAGAGTTAT	CTTCAACTAA	CTCTCCAATC	TCAAGAGCAG	АААТТТТАСТ	190200
CAAATCAGCC	AACTTTCTAA	TCTCACTGGC	CACAACAGCA	AATCCCTTTC	CCTCATCTCC	190260
TGCTCTTGCA	GCTTCAATAG	CCGCATTCAA	AGCAAGTAAA	TTGGTTTTTC	TAGCTATCTC	190320

			265			
TTCAATAACA	A CTAACTTTCT	CCACAATGTC		ATAACAGATT	CTTCAACGGC	190380
CCTACCACC	T ATCTGAGAA1	TTTCATTCGT	CTTTAAAGCT	' ATTTGTTCTG	TTTCATAAGA	190440
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ACTACTTATG	GGCAACATTG	САТААТААСА	ATCTTCTCCC	ATTTCGGACA	AAAGTATTCT	190980
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AAAAGGATTA	ACTGCTATAT	TGTTGGGATC	CACATAAATA	AAATTGCCTC	ТТТТАТАААА	191160
ACCGAATCTA	AATCTATCAA	AACTATCTGC	CACAATATCA	TTAAGCAAAT	ATCCGGCCAA	191220
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ATTGGTATTA	GCCTCAGAAT	GACCAAAATC	CATATTATTC	TCATGTCTTG	TGCTAACAAT	191460
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GGCCATAAAA	TTGTATAAGT	ATTGTCGATA	TTTTTTGCTC	ACCTTTACAG	AGTCAATAAC	191580
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CCTTGTAAGC	TGCTTATAAT	AATCTTCTAA	ATAACCGCAT	ААААСААААТ	TAAAAATCGT	191820
GGAAAAAAGT	AGCAGTATAA	AAATTAAAA	СААТААТААА	AATCCAACAA	ACCTGTATTT	191880
AAGCTTCAAT	AACATAATAA	ACTACCTCAC	AAATCACCTA	СТТАТТТААТ	САААТАААСТ	191940
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AAAAGAATTT	TCACAACATG	АААААСАААТ	ТАТАААСТАТ	ТАТТАТТААС	TGCTAATGCT	192060
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TTAGAATAAT	AATTAAATAT	GGGGTAATCA	CTAACATTTT	GGGAGGCATT	ATTAAAGAAA	192180
AAAAGGGTAA	TTGAGCCAAA	ACAATTGCCA	AAGTTTTCAC	AAATGAAAAC	AAAAAACTGC	192240
СТАТТААААС	TCCCAAAGGC	GTCCATTTTC	САААААТСАА	CATTACAATA	GCAATAAAAC	192300
CTTGTCCACC	TGTAACCCCT	TGCACATAAC	TTGATGCAAC	CACCGTTGTA	AGAACAGCAC	192360
CTGAAACCCC	TGCTAAAAAA	CCACTCAAAA	GAACGCAAAA	AAATCTAATT	TTATTTACAC	192420
TAACTCCAAC	AGACTCTAAT	ACCTCTGGAT	TTTCACCACT	TGCATTAATT	CTAAGCCCAA	192480
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AATATCTTTT	GCCAAAAATT	TGAAAAATAA	AAGATGTTTT	GTTTAAAATT	CCATCAAAAA	192600
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CAGCAAAAAT	TGAAAACAAT	GGATCATTTG	TAAAATATGC	AACTGTAGCT	CCTGAAAATG	192840
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GACCCCCAAG	ACCAGCTAAA	ATTAAGGTTT	GAGAATTTAT	TAAAGTTTCA	CTAATCAAGA	192960
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CCTTCTGGTG	AGAGCTTCCA	AGAAGCTAAA	AAATCAATAT	ATGCGCTTTC	TTTAATGGGT	193560
TTTGAAAAAT	CACTATTATC	TCTTTTAATA	AAACTAAAAT	СТААААТТАТ	AAATTTATTA	193620
TGAAATAATA	TCCAATTAAA	CATTATTCCT	GAAATCACTT	CGCTAATATT	GAATTTGGCT	193680
TTTAAATATC	CGATTAAAAT	TCCTAAACTG	CCTGAAGCTA	AAAAAGTAAT	ААТААААТА	193740
GTAATTACAT	GTAAAATTGG	AGGCAAATCA	AGTAAAACTG	ATGCTATTAA	AGCAACAATA	193800
GATCCTAGTA	TAAACTGGCC	TTCAACCCCA	АТАТТААААА	GACCCGCTTT	TAAAGAAATA	193860



			267			
CCAATAGAAA	GACCTGTAAA	AATCAAAGGA	GCTGAATAAC	ТТААААСАТА	ACCTAAATGT	193920
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ATTTAGCTTA	AGCCTATCAT	' CATTTTACCA	ATAACATCAA	TATCAAAATT	АТССТСТААА	194160
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TAGTTCGCTT	GAATCCTTAA	AGTTTATTTC	TTGACCTTTT	ААААТААТТС	GACCACTATT	195540
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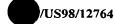


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GCAAATTAAT GT	ТААААТА	AAGAATAAAT	AAATTATTAC	AAAAGAGAGT	ATTATGAAAA	195960
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			271		_	
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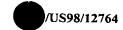
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AATCATAAAA	TTCTCTTGAA	AGATTTACAT	AATAAACTCC	ATTCTTTAAA	AAAGAATATA	205440
AAAATCTTGA	ATCACTTAAT	AAAAACCCAA	AAGAAAACCC	TTCATTGCTT	CCTAAAAGAA	205500
AATCTTTTAC	TAAAAGATCT	AAATTATCTT	TCAAATTTTG	TTCATCTCTT	AAATATCTTA	205560
AATTAGCAAC	AAATCCCTTG	CTAGAATGAA	AATAAAAAAC	CTTTTTTGAA	AAAAGATTAT	205620
САТААТТТАА	AAAAACCATA	CAAATAGAAA	ACAAAAAGCT	TACTGCCAAA	GACCCCAAAA	205680
GCACCCTAAT	CATATGCTCT	TTTTTTGAAT	ТТАААААТТТ	ATAAATCACT	ACATTATATT	205740
TAAAAAATAT	ATCCGAAATA	TTATTTTCA	TAAAAATTTA	TAAATTCCAT	TAAAGCTTTA	205800
AGTATTAGAA	TATTAAACTT	GCTCATGTAA	TTATAATCCA	AAATTAATTT	AGCATCTAAA	205860
ATATTGGATA	AAAACCCCAT	TTCAATCAAC	ACAGCAGGCA	TACTGCTGTT	TTTTATTACA	205920
AACCATTGCT	CTTTTCTGAT	TGGCCTAATA	TTAGTTTCGC	TTAACTCATT	TTTAAACACT	205980
ТТАТАСАААА	TTTCAGCCAA	TCTTTTTGAT	TCATATTTAT	ATTTAATATC	TAGTATATCA	206040
TTAAGCTCGC	TTAAGTATCT	ATTACCTTTA	ATATCATATC	ССТТААААТС	ТТТААТААСТ	206100
TCTCTTTTTG	AATCCTTAGG	AAGATACCAA	AACTCAACTC	CTCTAGCTTC	ACCGTTTGGA	206160
GCATCATTAG	CATGTATAGA	ТАААААТАТА	ACATTATTGG	GGAAATTTGG	CTTTATTGCA	206220
TTTGCAAATT	CCGACCGTTC	TTTTAAAGTT	АААТАААСАТ	CATTTATACG	AGTTAACAAA	206280



ATATTTTTAT TTA	CAAAATA ATTACTTAA	A ATTTTAGACA	AATATATAGA	ATAGGTTAAT	206340
GCAAAATCTT TTT	CCTGAAG CACAACGTC	А ТААССАТТТА	TCTTTAAAGT	CACAACAGCA	206400
CCAGTATCAT GCC	CGCCATG TCCAGGATC	A ATGATTATTG	AAGTAATTCT	GGGTTTATTA	206460
TAGTCTTTAA GAG	AACTAAA ATAATTTTC	A ATTTGTTTTA	ATACCTTTTG	ACTTATT <u>AA</u> A	206520
ATTTCTCCAC GAA	TGTCAAT AATTGGGTC	г асааасатат	AATAACCAGA	AGATGTAAGC	206580
GCATATTCAA AGC	CTACCCT AAACTTCAA	А ТАТСССТТАТ	CATTTTCAAT	TGTAAAAACA	206640
TCATTTTCAA TGT	TAAAATC AAACCTAAA	A ACATTAGTAT	САААААААТС	AAGAACATTT	206700
AAATAATCGG GGG	TCTTAGA ATACAAGCT	Г АААТАТGААА	ACAAAATCAA	ATCAATCAAT	206760
AATATCATTT TCC	CAAAGCT CAATGGCAC	r cttaagtct	CCTTTGAATC	TTAAGCTATT	206820
TTTAGTAATC TCA	CCTTTTA TCTCCTTAA	A CTCTTTTTGG	AAGAGAAAAG	GATTAATTTT	206880
GTATTTTAAA CAA	ATTTTGC AAAACCTTA	A AAAAGAAAAA	ACAACGCCAT	TTCCAACAAA	206940
AATTGGGACA TAA	ТТСТТАА ССАААААА	CAAAAACTT	TTGCTTTTGG	TTAAATTTTC	207000
TGGCGAATTT AATG	GCAGTAT ACTGGATTCT	CAGCTTTCTA	ТАААТАТААА	CATGCTCCCC	207060
AAAATAAATA GCTO	CTTAGTC CAAAATCTAT	TCTTTGAAAA	TATTCATTTT	GAATTCTTGC	207120
GTCAAAACCT CCAA	AGCTGTA AAAATTTCTC	TTTAGAATAA	AGTCCGCAAT	AATCCATGGT	207180
AATCAAAGTT TTTT	PCATAGT CTTTCTCAGA	ATTTACTAAA	ATTACCTTAA	ACTTTTGCTT	207240
TTTATCTATG CTG	GGAAGAA AAATTGAAGG	AATCATTTCC	TCTTCTTTAT	СААААААСТС	207300
CCCACCAACA AGAA	AGAACAT TTTTTTTAC	TATTTCATCG	AATATATTTG	GAATCCAAAA	207360
GGGATTTAAC AAGT	FACATAT CACTTTGCAA	AACAAAAACA	AAATCACAGC	TGGATTCTTT	207420
CATTGCTAAA TTAA	ACCTTTT CCCCAGAATT	' CAAATCGTCA	GAAAGTAAAA	ТАААТТТТАА	207480
CTTACCATAA CTTT	rctgaaa taaactgcaa	AGAACTTCTA	TTGCTCTGTT	TTTCAATTGA	207540
AATTATTTCT CTT	ATAAAGT CAAAATTTGA	ТАААААТТСА	AACAAATCTT	СТСТАААААТ	207600
TTTTGTTCCT CTGC	CTTAATA TTACAAAAGA	AATTCCAAAA	GAAGATTTTT	GTGAATAATT	207660
ATTTTTAGAT TGAA	ATAACAG TATATGAATA	GCCACTACCT	GGAAGACGCA	ТАААТТАСТТ	207720
TTAAAATCCT TATA	TAATAATATT AAATTAA	CATATGTTAC	АТААТАСААТ	GCTAATTGCA	207780
AGAATAATGA ATAT	ТТААТАС АТТАТТСТАС	GGCATGATCA	ТТАТСАТТТТ	TGCACTCATT	207840
	AGAATAT ACAGTACGAC				207900
AAAATTGAAT ATAA	AAATAGA CTCAGAAAAT	GACTTTATAG	CATTTAAAGA	ТАТАААСААТ	207960
AACGAAAAAG AAGA	AGTAAT CATCAGATCA	AGACTAAACT	CATATAAAAA	TTCAAAGATA	208020
					•



AGAGAAARAT TIGGAATTGT TAAAGTATTT GATATAAACA CACCARAART AAAAGAAATA 208080 TCTGACTCGC TTATGAGGGA TAGTTATAAT AACAGAGTAT TIGGATCGTG GAGATTATT 208140 CATAATGCAG AAAGAGGAAT CAACTCTTTG GTATATATTG TAAAAGCAGA AGAATTTGCA 208200 AATGATACAT TITTGCTTGA TGCAATTGAT GAGATTGCT CAACAATAAG TATTTTCAAA 208260 AAAATAATAA CAACCAACAA CGAAAACATT GATAATAATG AAGAAAATAA CAATACAAAT 208320 GAATCAAATG AACAGCCCCC CTTAAAGCAA GAAAACAA ATTCAACAAA AGAATCTAAT 208380 AACGAACTTA AAGAAGATCA AATAGAAGAA GAACTTCAAG AAATGAACCTT GTCACCTATT 208440 CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCACAA AATGAACCTT GCCCCTATT 208560 TTTATTTGAA ATTTGCCCT AGAGTACGC TGAAGCTCAA GAGCATACCT TACCTTATAA 208620 CTTGAATCAA TATAAGCTAT TCCAAGGGC TCTAAATCAT AAATCTTT TTGAGCACC 208740 CTTGAATCAA TATAAGCTAT TCCAAGCAGC AAAGGTGTAT TTCCATCCA AAAAGCAAAA 208740 ATTATACATAA TATTTTACCCCTA AAGACCATT CCATTGCCAT ATTCAACATT TTGAGCACCA 208800 TTTTACATAA TTTCTTTAC GTAAAAATGA TCCATTATT TAAAAAAAGA CATCAACAAA 208920 ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATA AATTTTTATT TTAAAACAAA 209920 ATTATATATT GAAATATATAA ATTTTTATAAATTTA AATTTTTTAT TAAAACAAAA 209100 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGGTATT ACTAGGTTAT CAACTTCTT CAACACACAAA 209100 AGTGATAAAT			2/5			
CATAATGCAG AAAGAGGAAT CAACTCTTTG GTATATATTG TAAAAGCAGA AGAATTTGCA 208200 AATGATACAT TITTGCTTGA TGCAATTGAT GAGATTGCT CAACAATAAG TATTTTCAAA 208320 AAAATAATAA CAACCAACAA CGAAAACATT GATAATAATG AAGAAAATAA CAATACAAAT 208320 GAATCAAATG AACAGCCCAC CTTAAAGCAA GAAATAACAA ATTCAACAAA AGAATCTAAT 208380 AACGAACTTA AAGAAGATCA AATAGAAGAA GAAATACAAA ATTCAACAAA AGAATCTATT 208440 CAAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGAACCTT GTCACCTATT 208500 TTTATTTGAA ATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCT TACCTATAAA 208620 CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA 208680 TTTTGATCTT TTTTAAAACA AAAAAGCATT CCATTGCCAT ATCAACTTT TTGAGCACCC 208740 ATTATCATAA TTTCTTTAC GTAAAAATGA TCACCAAAAG ATAAAAAAAA CTTAACCCCA 208800 ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATT 208980 ACTATTTTTTTTT GAAAATTATT TACATTATA ATTTTTTAT TAAAAACAAA 209100 ACTATTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT						208080
AAATAATAA TATTOCTTGA TGCAATTGAT GAGATTGCCT CAACAATAAG TATTTTCAAA 208320 GAATCAAATA CAACCAACAA CGAAAACATT GATAATAATG AAGAAATAA CAATACAAAT 208320 GAATCAAATG AACAGCCCAC CTTAAAGCAA GAAAAAACAA ATTCAACAAA AGAATCTAAT 208380 AACGAACTTA AAGAAGATCA AATAGAAGAA GAACTTCAAG AAATCAAAG CCAATAATTT 208440 CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGCACTT GTCACCTATT 208500 TTTATTTGAA ATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCT TACCTTATAA 208560 AGAGAATTAA CATTTGCCCT AGAGTACGC TCTAAATCAT AAATCTCTT AATAATTCCA 208620 CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTCCATCCA AAAAGACAAA 208680 TTTTGATCTT TTTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC 208800 TTTATCATAAA TTTCTTTATC GTAAAAATGA TCAGCAAAAA ATAATTTTTT TTAAAACAAA 208860 ACTAAAAACA AAAACCGTTT TAAAATTTT TTCAATTAAC ACCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTTAT TTAAAACAAT 208980 AATAGAAACA AAAACCGTTT TAAAATTTT TACAATAAA ATATTTTTAT TTAAAACAAT 2099040 CTTTTTCTTT ACCTCAATAA AAATGCATA CTGAGAATAT AATTTTTTAT TAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATTATTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGAAAAT TCAACGTCTT CAACAACAAA 209220 ATAAAAAAAT GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATCCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA ACCCTTATGCC 209340 ATAAAAAAAT GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATCCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA ACCCTTAGCC 209340 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG ACCCCAGAAC 209280 AAAAGATCCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA ACCCTTAGCC 209340 ATAAAAAAATA TAAAGAAAAA GTATAACAAA TAAAAAAACTT GAAAAACATA AAGGAAAAT CAAAAACAAA CCCTTTAGCC 209340 ATAAAAAATA TAAAGAAAAA GTATAACAAA TAAAAAAACTT GAAAAACAT GAAAAACAAA 209000 ATCAATTATA ATCCCAAAAT GGAAAACCCT TACAAAGGAC TTAAAGAAC GCCTTATCACT 209460 TGCACTTCTT GCCTTTCTT TTCTTTTAAA AACTATTA TAAAAAAAAT AAAGGAAAAT 209500 TGGACATTCG CTCTTTCCCAAA AGCAGAAAC TTCAAAATTT AAAAAAAAAT AAAGGAAAATT 209560 TGGACATTCG ATATCACCAA GCTTACACAA ACGAAAACT TAAAAAAAATT AAAAAA						208140
AAAATAATAA CAACCAACAA CGAAAACATT GATAATAATG AAGAAAATAA CAATACAAAT GAATCAAATG AACAGCCCAC CTTAAAGCAA GAAAAAACAA ATTCAACAAA AGAATCTAAT 208380 AACGAACTTA AAGAAGATCA AATAGAAGAA GAAAAAACAA ATTCAACAAA AGAATCTAAT 208440 CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGAACCTT GTCACCTATT 208500 TTTATTTGAA ATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCTT GCCCCTATT 208560 AGAGAATTAA CATTTGCCCT AGAGTACGGC TCTAAATCAT AAATCCTTT AATAATTCCA 208620 CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAAGGTGTAT TTCCATCCA AAAAGACAAA 208680 TTTTGATCTT TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC 208800 TTTATCATAAA TTTCTTTATC GTAAAAATGA TCAGCAAAAG ATAAAAAAAA CTTAACCCCA AGCAAAAAACA AAAACCGTTT TAAAATTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAAATTT GAAAATCAAG ATTTTAAAGT AATTCTTAAAA ATATTTTTAT TTAAAACAAT AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGGTGTAG CCTCGCCAAA AGGGTTACGAA AACACTTTTAT TTTTTCTCAAA ATATTTTTAT CAAAAACAAA AGGGAATAAAA TGCACAAAAA AAAATCGATA CTGAGAAAAG GCATATTTA AAATCATTT ACCTCAAATAA AAAATCGATA CTGAGAAAAT TAACTTTTTAT CAAAAACAAA AGGGAATAAAAT TGTAAAGTCT GCTTCCTAGC ATAAAGAAAT TCAACTTCTT CAACAATCATA AGGCAATAAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACAACCAAA ATAAAAAAATA TGTAAAAGTA ATAAATAAAA AACTGTAATA CCCATACCAAA GCCTTTAGCC AAAAGAATAA TAAAGAAAAA ATAAATAAAA AACTGTAATA CCCATACCAAA GCCTTTAGCC AAAAGAATAA TAAAGAAAAA ATAAATAAAA AACTGTAATA CCCATACCAAA GCCTTTAGCC 209340 ATAAAAAAATA TAAAGAAAAAA TAAAAAACAT GAAAATAAAAT	CATAATGCAG AAAGAGGAA	T CAACTCTTTG	GTATATATTC	TAAAAGCAGA	AGAATTTGCA	208200
AACGAACTTA AACAGCCCAC CTTAAAGCAA GAAAAAACAA ATTCAACAAA AGAATCTAAT AACGAACTTA AAGAAGATCA AATAGAAGAA GAACTTCAAG AAATCAAAGC CCAATAATTT CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGAACCTT GTCACCTATT CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGAACCTT GCCACCTATT TTTATTTGAA ATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCTT TACCTTATAA AGAGGAACTTAA CATTTGCCCT AGAGTACGGC TCTAAATCAT AAATCCTTT AATAATTCCA AGAGAATTAA CATTTGCCCT AGAGTACGGC TCTAAATCAT AAATCCTTT AATAATTCCA CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA CTTTGAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA ATGTAACCTT TTGCCCTATC CAAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA ACTAAAAACA AAAACCGTTT TAAAATTTT TTCAATTATC AGCCTTATTA AAAATCATTT AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGCGGTT ACTAGTGTAG CCTCGCCAAA AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA AGTGTACGAA AACATTTTAT TTTTGCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT AGCAATAAAAT TGTAAAGCTC GCTTCCTAGC ATAAAGAAAA CACCAGAAC 209200 AGCAATAAAAT GTGACTTGA GAACATTCC TTTAACAGAA TCCATCCTT CAACATCATA AGAAAAAAAATA TGTAAAACAC GCATACTATA CCCATACAAA GCCTTTAGCC ATAAAAAAAAA GTGCTCTTGA GAACATTCC TTTAACAGAA TCCATGCTAG AACCCAGAAC AAAAGAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAA AAAGCCAAAAT ATAAAAAAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAA AAAGCCAAGAAC ATAAAAAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAA AAAGCCAAGAAC ATAAAAAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAA AAAGCCAAGAAC ATAAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAA AAAGCCAAGACC CTCTATCACT ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAAGGTC TTAGAGCTC CTCTATCACT COTGAATTTT TACTTCTTT TTCTTTTAAA GGCCTTTCTT TTAAAATATT TACAAAACTT GAAAATAAAT CCCATCAATT COGGACATTCG CTCTTTCTT TTCTTTTAAA AGCAAAACTT GAAAATAAAT AAAGCAAAATT COGGACATTCG CCACATTGCT CGCAAAAAAC TTCAAAAATTA TACTTTCCCA AACGAAAATT COCTCAAATTT TACATAAAA TAAGGAATCT TAAAAACTT TACTTTCCCA AACGAAAATT COCTCAAATTT TACATAATATT AAAGAATATTA AAATCATTA TAATTAAAAAA COOTAGATT TACA	AATGATACAT TTTTGCTTG	A TGCAATTGAT	GAGATTGCCT	CAACAATAAG	TATTTTCAAA	208260
AACGAACTTA AAGAAGATCA AATAGAAGAA GAACTTCAAG AAATCAAAGC CCAATAATTT CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGAACCTT GTCACCTATT 208500 TTTATTTGAA ATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCT TACCTTATAA 208560 AGAGAATTAA CATTTGCCCT AGAGTACGGC TCTAAATCAT AAATCTCTT AATAATCCA 208620 CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA 208680 TTTTGATCTT TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTCAATAAAA CTTAACCCCA ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTCAATAAAA CTTAACCCCA 208800 TTTATCATAA TTTCTTTATC GTAAAAATGAT TCAACAAAGA ATAAAAAAGC CATCGACAAA ACTAAAAACA AAAACCGTTT TAAAAATTTT TTCAATTATC AGCCTTATTA AAAATCAATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATTTTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209100 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA AGCAATAAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA ATAAAAAATA GTGCTCTTGA GAACATCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC AAAAGAATA TAAAGAAAAA GTATAAAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC ATAAAAAAATA TAAAGAAAAA GTATAAAAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAACTGAT GCCTACAATT TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT CTGGACATTCG ATATCACCAA GCTTAGCAG CAGGTGGGA CAAACAGATC GACTACATT COGACATTCG ATATCACCAA GCTTAGCAG CAGGTTGGGA CAAACAGATC GATTAAGTCC AACCTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT CCCTCAAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAAATA TAATAAAAAA 209760 CCCTCAAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAA 209760	ААААТААТАА СААССААСА	A CGAAAACATT	GATAATAATG	AAGAAAATAA	CAATACAAAT	208320
CAAAATCATT CTACTAATAA AGAATTAACA TCAAAGCAAA AATGAACCTT GTCACCTATT 208500 TTTATTTGAA ATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCT TACCTTATAA 208560 AGAGAATTAA CATTTGCCCT AGAGTACGGC TCTAAATCAT AAATCTCTTT AATAATTCCA 208620 CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA 208680 TTTTGATCTT TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC 208740 ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CATCAACCCA 208860 ACTAAAAACA AAAACCGTTT TAAAAATTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTTAT CAAAAACAAA 209100 AGCAATAAAAT TGTAAAGTC GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCT TTTAACAGAA TCCATGCTGA AACCCAGAAC AAAAGAATAA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAAAAAATA GTGCTCTTGA GAACATCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC ATAAACAATA TAAAGAAAAA GTATAAAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAAAAATA TAAAGAAAAA GTATAAAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAAAGTCC TCCTACCT TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGACTTTCT TTAAAGTACT GCAAAACAAAT 209520 GTAGTGACAA TTAGGACAAC CTTCCCAAAA GCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAG ACAGTTGGGA CAAACAGATC GATTAAGTCC 209540 AACCTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT AACTTAATA TAATAAAAAA 209700 CCTCAAAATTT TACTTAAAAA TAAGTATTTA AAAGAAAATT TAATTAAAAAA 209700 CCTCAAAATTT TACTTAAAAA TAAGGAATCTTA AAATACTATA TAATTAAATAA TAATAAAAAA 209700 CCTCAAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAAATAA TAATAAAAAA 209760	GAATCAAATG AACAGCCCA	C CTTAAAGCAA	GAAAAAACAA	ATTCAACAAA	AGAATCTAAT	208380
AGAGAATTAA CATTTAGAAAA TGAGCCCCTT GGAAGCTCAA GAGCATACCT TACCTTATAA 208600 AGAGAATTAA CATTTGCCCT AGAGTACGC TCTAAATCAT AAATCTCTTT AATAATTCCA 208600 CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA 2086800 TTTTGATCTT TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC 2087400 ATGTAACCTT TTGCCCTATC AAGCCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA 208800 ATGTAACACTT TTGCCCTATC AGGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA 208800 ACTAAAAAACA AAAACCGTTT TAAAAATTTT TCAAGTATAC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAATTT TAAAAACAAT 2089800 ACTAAAAAACA AAAACCGTTT TAAAATTTT TAAAATTATC AGCCTTATTA AAAATCATTT 2089800 ACTATATAATTT GAAATATAAG ATTTTAAAAT AATTTTTATT TTAAAAACAAT 2099040 ACTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGGATATT AATTTTTATT CAAAAAACAAA 2091000 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAAACA GCATATTTTA AATGGGTTTT 2091600 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGAAAAC GCATATTTTA AATGGGTTTT AAAAAAAAAA	AACGAACTTA AAGAAGATC	A AATAGAAGAA	GAACTTCAAG	AAATCAAAGC	ССААТААТТТ	208440
CTTGAATCAA TATAAGCTAT TTCAAGCAGC TCTAAATCAT AAATCTCTTT AATAATTCCA CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA CTTGAACCTT TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA CTTATCATAA TTTCTTTATC GTAAAAATGA TCAGCAAAAG ATAAAAAAGA CATCGACAAA CCTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATTT CAAAAACA AAAACCGTTT TAAAATTTTT TCAATTATC AGCCTTATTA AAAATCATTT CAAAGAACA AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGAGAAT TCAACTTCTT CAACATCATA AGCAATAAAT GTGCTCTTGA GAACATTCT TTTAACAGAA TCCATGCTAG AACCCAGAAC AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC ATAAAAAAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC ATAAAAAAAA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACCTT GAAAATAAATA AAAGCAAAAT ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACCTT TTAAGCTCTT GAAAAGGACT CTCATACCAT GCCTACCAAA GCCTACCAAA GGCCTTTCCTT TAACCATCTT GCCTACCATT COOPSEO GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAAT GCCTACAATT COOPSEO GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT COOPSEO CCTCAAAATTT AACCTAATTT AAGGAATACTT AAAAAACTTA TAATTAAATTA	CAAAATCATT CTACTAATA	A AGAATTAACA	TCAAAGCAAA	AATGAACCTT	GTCACCTATT	208500
CTTGAATCAA TATAAGCTAT TTCAAGCAGC AAAGGTGTAT TTTCCATCCA AAAAGACAAA 208800 TTTTGATCTT TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC 208740 ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA 208800 TTTATCATAA TTTCTTTATC GTAAAAATGA TCAGCAAAAG ATAAAAAAGA CATCGACAAA 208860 ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTTAT CAAAAACAAA 209100 AGGGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCT TTTAACAGAA TCCATGCTAG AACCCAGAAC 209340 ATAAACAATA TAAAGAAAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GAACATACAT GAAAAGAAT 209580 TGGACATTCG CTCTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCCCAAA AGCAGAAACT GGACCTACAAT 209580 TGGACATTCG ATATCACCAA GCTTAGCAG ACAGTTCGGA CAAACAGATC GATTAAGTCC 209540 AACTTTTTCG CCACATTGCT CGCAAAAAAC TACAAATTT ACCTTTCCCA AACGAAAATT 209500 CCTCAAAATTT TACCTAAAAA TAAGTATTA AAAAAACTT AAATAAATAA AACTTTCCCA AACGAAAATT 209700 CCTCAAAATTT TACCTAAAAA TAAGTATTTA AAAAAACTT TAATTAATTA TAATAAAAAAA 209760	TTTATTTGAA ATTTAGAAA	A TGAGCCCCTT	GGAAGCTCAA	GAGCATACCT	ТАССТТАТАА	208560
ATGRANCET TTTTAAAAAC AAAAAGCATT CCATTGCCAT ATTCAACTTT TTGAGCACCC 208800 ATGRAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA 208800 TTTATCATAA TTTCTTTATC GTAAAAATGA TCAGCAAAAG ATAAAAAAGA CATCGACAAA 208860 ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAA AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAGAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACTTAAAAA TAAGGAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACTTAAAAA TAAGGTATTA AAAAAAACTT AAATTAATTA TAATAAAAAAA 209760	AGAGAATTAA CATTTGCCC	P AGAGTACGGC	TCTAAATCAT	AAATCTCTTT	ААТААТТССА	208620
ATGTAACCTT TTGCCCTATC AAGCTCATTA GATGCTATTT TTACAAAAAA CTTAACCCCA 208860 TTTATCATAA TTTCTTTATC GTAAAAATGA TCAGCAAAAG ATAAAAAAG CATCGACAAA 208860 ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAG ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACTTAAAAA TAAGGTATTA AAAATACTTA TAATTAATTA TAATAAAAAAA 209760	CTTGAATCAA TATAAGCTA	TTCAAGCAGC	AAAGGTGTAT	TTTCCATCCA	AAAAGACAAA	208680
TTTATCATAA TTTCTTTATC GTAAAAATGA TCAGCAAAAG ATAAAAAAGA CATCGACAAA 208860 ACTAAAAACA AAAACCGTTT TAAAATTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209340 AAAAGATGCA AAAATGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATT TACTTAAAAA TAAGTATTT AAAGAAAAAC TTCAAAATTT TACTTAAAAAA TAAGTATTT AAAGAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATT TACTTAAAAA TAAGTATTT AAAGAATATA TAATTAAATAA TAAATAAA	TTTTGATCTT TTTTAAAAA	AAAAAGCATT	CCATTGCCAT	ATTCAACTTT	TTGAGCACCC	208740
ACTAAAAACA AAAACCGTTT TAAAATTTTT TTCAATTATC AGCCTTATTA AAAATCATTT 208920 ATTATAATTT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAACT GGACCTACAAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATT TACCTAAAAAA TAAGTATTTA AAAAAAACT TAAATAAATA TAAATAAAAAA TAAATATTA AACGAAAAA TAAGTATTTA AAAAAAACT TAAATAAATA TAAATAAAAAA TAAAACAATT TACCTAAAAAA TAAGTATTTA AAAAAAACT TAAATAAATA TAAATAAAAAA TAAAACAATT TACCTAAAAAA TAAGTATTTA AAAAAAACT TAAATAAATA TAAATAAAAAA CCCTCAAAATTT TACCTTAAAAAA TAAGTATTTA AAAAAAACTT TAAATAAAAAAA TAAAAAAAAATTTA AAAGAAAAAA TAAAAAAACTT AAATAAATTA TAAATAAAAAAA CCCTCAAAATTT TACCATAAAAA TAAGTATTTA AAAAAAACTTA TAAATAAATTA TAAATAAAAAAAA	ATGTAACCTT TTGCCCTATO	AAGCTCATTA	GATGCTATTT	ТТАСААААА	CTTAACCCCA	208800
ATTATAATT GAAATATAAG ATTTTAAAGT AATTCTTAAA ATATTTTAT TTAAAACAAT 208980 AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATT TACATAATATT AAGGAATACTT AAAGAATAT TAATTAAAAAAA 209760	TTTATCATAA TTTCTTTATC	GTAAAAATGA	TCAGCAAAAG	ATAAAAAAGA	CATCGACAAA	208860
AATAGAATCA CCAAGATCCC ATAAATAATT CAAAGGCGTT ACTAGTGTAG CCTCGCCAAA 209040 CTTTTTCTTT ACCTCAATAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACTTAAAAA TAAGTATTTA AAAGTAATTA TAAATAAAAAA 209760	ACTAAAAACA AAAACCGTTT	TTTTAAAAT	TTCAATTATC	AGCCTTATTA	AAAATCATTT	208920
AGTOTACGAA AAATTGCATA CTGAGATATT AATTTTTAT CAAAAACAAA 209100 AGTGTACGAA AACATTTTAT TTTTGTCAAA ATTAAAAACA GCATATTTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACTTAAAAAA TAAGTATTTA AAAGAATATA TAATTAAATAA TAATAAAAAAA 209760	ATTATAATTT GAAATATAAG	ATTTTAAAGT	AATTCTTAAA	ATATTTTTAT	ТТААААСААТ	208980
AGTGTACGAA AACATTTAT TTTTGTCAAA ATTAAAAACA GCATATTTA AATGGGTTTT 209160 AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACCATATTT AAGGAATACT AAAACAGATA TAAATAAAAAA 209760	AATAGAATCA CCAAGATCCC	TTAAATAATT	CAAAGGCGTT	ACTAGTGTAG	CCTCGCCAAA	209040
AGCAATAAAT TGTAAAGTCT GCTTCCTAGC ATAAGGAAAT TCAACTTCTT CAACATCATA 209220 ATAAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAAACTT GAAAATAAAA AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAA 209760	CTTTTTCTTT ACCTCAATAA	AAATTGCATA	CTGAGATATT	AATTTTTTAT	САААААСААА	209100
ATAAAAAATA GTGCTCTTGA GAACATTCTC TTTAACAGAA TCCATGCTAG AACCCAGAAC 209280 AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAAA 209760	AGTGTACGAA AACATTTTAT	TTTTGTCAAA	ATTAAAAACA	GCATATTTTA	AATGGGTTTT	209160
AAAAGATGCA AAAATAGAAA ATAAATAAAA AACTGTAATA CCCATACAAA GCCTTTAGCC 209340 ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAA 209760	AGCAATAAAT TGTAAAGTCT	GCTTCCTAGC	ATAAGGAAAT	TCAACTTCTT	CAACATCATA	209220
ATAAACAATA TAAAGAAAAA GTATAACAAA TAAAAAACTT GAAAATAAAT AAAGCAAAAT 209400 ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAA 209760	ATAAAAAATA GTGCTCTTGA	GAACATTCTC	TTTAACAGAA	TCCATGCTAG	AACCCAGAAC	209280
ATCAATATTA ATCCCAAAAT GGAAAAGCCT TACAAAGGTC TTAGAAGCTG CTCTATCACT 209460 TACCATTCTT GCTCTTTCTT TTCTTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAAATTT TACTTAAAAAA TAAGTATTTA AAACTATTA TAATTAATTA TAATAAAAAAA 209760	AAAAGATGCA AAAATAGAAA	ATAAATAAAA	AACTGTAATA	CCCATACAAA	GCCTTTAGCC	209340
TACCATTCTT GCTCTTTCTT TTCTTTAAA GGGCTTTTCT TTAAGTTCTT GAAAAGGACT 209520 GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAAGTATTA TAATTAATTA TAATAAAAAA 209760	АТАААСААТА ТАААGААААА	GTATAACAAA	TAAAAAACTT	GAAAATAAAT	AAAGCAAAAT	209400
GTAGTGACAA TTAGGACAAC CTTCTCCAAA AGCAGAAACT GGACCTACAT GCCTACAATT 209580 TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAAGTATTA TAATTAATTA TAATAAAAAA 209760 TATATGAATA TTACATATTT AAGGAATACT AAAACATGAA ATGAATACTATA TAATAAAAAAA	АТСААТАТТА АТСССААААТ	GGAAAAGCCT	TACAAAGGTC	TTAGAAGCTG	СТСТАТСАСТ	209460
TGGACATTCG ATATCACCAA GCTTAGCAGC ACAGTTGGGA CAAACAGATC GATTAAGTCC 209640 AACTTTTTCG CCACATTGCT CGCAAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAACATATA TAATTAATTA TAATAAAAAA 209760 TATATGAATA TTACATATTT AAGGAATACT AAAACATGAA ATGAATA TAATAAAAAA	TACCATTCTT GCTCTTTCTT	ТТСТТТТААА	GGGCTTTTCT	TTAAGTTCTT	GAAAAGGACT	209520
AACTTTTCG CCACATTGCT CGCAAAAAC TTCAAAATTT ACCTTTGCCA AACGAAAATT 209700 CCTCAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAA 209760 TATATGAATA TTACATATTT AAGGAATACT AAAACATGAA ATTACATATA TAATAAAAAA	GTAGTGACAA TTAGGACAAC	CTTCTCCAAA	AGCAGAAACT	GGACCTACAT	GCCTACAATT	209580
CCTCAAATTT TACTTAAAAA TAAGTATTTA AAATACTATA TAATTAATTA TAATAAAAAA 209760	TGGACATTCG ATATCACCAA	GCTTAGCAGC	ACAGTTGGGA	CAAACAGATC	GATTAAGTCC	209640
TATATGAATA TTACATATTT AAGGAATACT AAAAGATGA ATTACAT	AACTTTTTCG CCACATTGCT	CGCAAAAAAC	ТТСААААТТТ	ACCTTTGCCA	AACGAAAATT	209700
TATATGAATA TTACATATTT AAGGAATACT AAAACATGAA ATCGGGATTT GCAGCAATAC 209820	ССТСАААТТТ ТАСТТААААА	TAAGTATTTA	AAATACTATA	ТААТТААТТА	AAAAAAA	209760
	TATATGAATA TTACATATTT	AAGGAATACT	AAAACATGAA	ATCGGGATTT (GCAGCAATAC	209820



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TTGGTAGACC	ATCAACTGGA	AAATCTACC	C TTTTAAATTO	C AATATGCGG	A CATAAAATAT	209880
СААТААТАТС	CCCTATTCCG	CAAACAACT	A GAAATAATAT	C AAAAGGAATO	TTTACGGACG	209940
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			AAAAGTAAGG			210960
			TATGCCAGAA			211020
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					TTGCACTTGA	211200
					TTGAAGAAAA	211260
TTGGTATCCA .	ATCAGTGTAT	TTTGTACAAA	ATGCAATAGA	GACACAACAA	CTGTAAATAA	211320
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			277			
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АААТТААТАТ	ACTTGGACAA	AGAAGTGAAA	ATAAATTTTA	AATTATCTTG	ССТАGАТААА	214920
САААААТСТА	CGTGCTTAAT	GTAAAGTGTT	TTTATAATTG	GATGACCCAG	CAAGATTAAA	214980
TGACATTTTA	ATTCCTCTAA	AATTCTATTA	AAAATGCTAA	AATCTTTACA	ACAAAATGAC	215040
АААТТААТАТ	AAAGACTTAA	CTCGACAAGA	GGTAAAAGAT	CTACTCTCAT	AATTGGTCTT	215100



			279			
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GCAGAAGCTT	TAATGCCAAG	ATCAATAGTC	CTAGCACCAA	GAGTAGGCCC	TATTATTCTC	218640

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			281			
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285	
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287		
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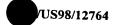
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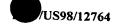
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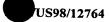
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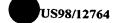
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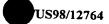
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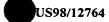
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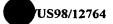


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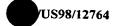
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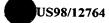
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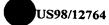
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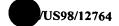


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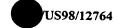
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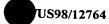


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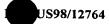
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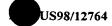


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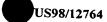


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AAAATGCTCA	AATAAAATTG	ATAATTGCAT	TTTACAATAA	САТААТТАТТ	TCTGGCATAA	306660
TAGTGGGAAT	TTACAAAGAA	AAAGCAGTCT	ATCTTTATGG	GGCTTCAAGC	AAGGAATATA	306720
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AAATAAAAGA	ATATGATTTA	TTAGGAATTC	CCCCAATTGC	АААТАААААС	CACCCCTTAT	306840
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ACTTCTATTA	CAAAGTAATA	AGGAAAAAA	TTTAACCAAG	ATTATTAACT	AAATTTTTAA	307020
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			331			
AATCAAATAG	AAAATATTGT	ATGCTACTTT	TTTCTAAAAT	CTTAATAATT	TTTACAGTTG	307200
CACTACCTCT	TATTAAAATC	CAAGTTCTCA	CAATATCTGC	AATCTTGCTA	АААСТТАААА	307260
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CTTTAGGAAT	ТАТАААААСА	GCTCGTGAAT	TAGAAAAACT	TCCAATCAAG	CTGTCATTAA	307560
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CGGAAAATAA	AATAACTCTG	ACTTTTGATT	TAAATTTTGA	AAAATATTTG	CAATAAGCCT	307680
GATCCTGAAT	AATCACAATT	CCTGAAGTTT	GATTTACAAA	AATTTTTGAC	ТТАТСААТАА	307740
TATAATCATC	AAAATTTAAA	TAATAGTTTT	GATGATCGTT	GTAAACATTT	GTAATAATAC	307800
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САААТАТАТТ	GCAACTTTCA	TGATTTCTTT	TTTACTATTA	TGCATTTCAT	AGCCCATCAT	308820
AATATTTAAA	GAAATACAAA	AGGAATGAAT	TATATTAAAG	ТАААТААТА	AAATAGAAAA	308880
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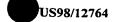
				•		
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TGCAATAGCA	GCTCCAACTA	CTCCCATATG	AAAAACAAAA	АТААААТАТ	AATTAAATAC	309180
AAAATTCAAA	AGCACTGAAA	AAACAGAAAT	GTAAACTTGA	AATTTAACAA	TTTCAACAAC	309240
CTTAAGAGCA	TTAGCAATAA	GTCCTTTTAT	GATTGCAAAT	ACGAAAGAAA	AAATAGCTAT	309300
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TAAAAGATCT	ТТАТАААТТТ	TATCCTTTTT	CGATACACCC	AATGAATACA	TAAATTTCTC	309720
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CTCTCGTATC	CAATCATCGT	GAAAAAATTC	ААТТАТААТА	AATATATTT	TAATAATCTC	309960
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AGGATATAA TTAAAAACAA TATTTATTAA AACAACAATA GAAGTAACAT ATAAGGGTAT 310740 TTTTACTTCT TTAGCACTCT TAAATCCCAT AGCAGATAA AAAGAATATG CCATTAGCAC 310800 ATAAGACAAT GAAATAATTT TTAAATACTC TGATCCAAAA TTTAAAGAAT CCTGATTAGCAC 310860 TGTAAATAGC TTATAATGT TCTTGGAAA AAGAATGAA AATATAAAAA AAATTATCC 310920 TATTGTAGTT CCAATTAATA ATATATAAGC AAAGAATGAA CATAGCAAA AAAATATT 310980 TTTTGCAATT GCTGGGAAA CGTAAGGGCT TAAAGCCGTA CCAAGTCCAA ACACAATAAT 311040 AAAAAAAGA AAAGTTACCC TATTTGCCAA AGAAACTCCC GTAACACTGAA AAGAGCCCAA 311100 ATAGGAGATC ATAAAATATT CAAAAAAAGT TACCATTTGA AATAAAAGCG ATTCAAACTTTT 311220 GATCATATTC CTTATATACA GCCCTTT ATTATTACTG TCATAATTTT TAAACTTTTT 311280 CTAGAATATA CATATATTA GAATTAACTA TTAATTACT TAAATTATAT TATAATTAAA 311400 TAAATATGAAA TCAATTATTA GAATTAACTA TTAATTATAC TAAATTATAT TATAATTAAA 311400 CAACATTTCA AAAAAAAATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311520 AAGACGAAAAA TTATTTTAGAA AAAATCCTAT TCAGTCTTA AAAGAAATAA AACCCTTAGCT 311580 TTATAAAATA CCTGATCTGG TAAATATTGA TGATTTTTGAA AAATTAAAA AAAATACCAA TTCCTGAAAA 311700 AAAAAAAATTAT GGGATTGAAA TTAAAATATA TGACGACTAA ATTCGACCAAAAATTA TACCACACAAAAA AAATATTAAA AATTCGACCAAAAAATAA AAATATCAAA ACCCCACAACAAAAAAAA			333			
ATAMGACAAT GAAATAATTT TTAAATACTC TGATCCAAAA TTTAAAGAAT CCTGATTAGC 310920 TGTAAATAGC TTATAATGT TTCTTGGAAA AAGAAATGAA AATATAAAAA AAATTATTCC 310920 TATTGTAGTT CCAATTAATA ATATATAAGC AAAAGAATGT CTTACTGAA GAAACTTCTT 310980 TTTTGCAATT GCTTGGGAAA CCTAAGCGCT TAAAGCCGTA CCAAGTCCAA ACACAATAAT 311040 AAAAAAAGA AAAGTTACTC TATTTGCCAA AGAAACTCCC GTAACATGAA AAGAGCCCAA 311100 ATAGGAGATC ATAAAATTAT CAAAAAAAGT TACCATTTGA AATAAAAGCG ATTCAAAAGC 311160 TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTGG TCATAATTT TAAACTTTT 311280 CTAGAATATA CATATATTA GAATTAACTA TTAATTACC TAAAATATAT TATAATTAAAA 311400 CAACATTCC CTTATATACA GTCTCCTTTA TTTATTATCT TAAATTAAT TATTAATTA						310740
TOTAMATAGE TITATAATGT TICTTOGAAA AAGAAATGAA AAATATAAAA AAATTATTCC 310920 TATTGTAGTT CCAATTAATA ATATATAAGE AAAAGATTGT CTTACTTGAA GAAACTTCTT 310980 TTTTGCAATT GCTTGGGAAA CGTAAGCGCT TAAAGCCGTA CCAAGTCCAA ACACAATAAT 311040 AAAAAAAGA AAAGTTACTC TATTTGCCAA AGAAACTCCC GTAACATGAA AGAGCCCAA 311100 ATAGGAGATC ATAAAATTAT CAAAAAAAGT TACCATTTGA AATAAAAGGG ATTCAAAAGC 311160 TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTG TCATAATTTT TAAACTTTTT 311220 GATCATATC CTTATATACA GTCTCCTTTA TTTATTACTT TTAACATAAT TTAAACTAAA 311340 CTAGAATATA CATATATTA GAATTAACTA TTAATTATCT TAAAATTAAT TATAACTTAAAA 311400 CAACATTCA AAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311400 CAACATTCA AAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311400 AAGACATATCA AAAAAAAAAAAAAAAACC TCTTAACTCT AATAAATA						310800
TATTGTAGTT CCAATTAATA ATATATAAGC AAAAGATTGT CTTACTTGAA GAACTTCTT TTTTGCAATT GCTTGGGAAA CGTAAGCGCT TAAAGCCGTA CCAAGTCCAA ACACAATAAT 311040 AAAAAAAAGA AAAGTTACTC TATTTGCCAA AGAAACTCCC GTAACATGAA AAGAGCCCAA 311100 ATAGGAGATC ATAAAATTAT CAAAAAAAGT TACCATTTGA AATAAAAGGG ATTCAAAAGC 311160 TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTGG TCATAATTTT TAAACTTTTT 311220 GATCATATTC CTTATATACA GTCTCCTTTA TTTATTACTT TTAACTAAAT TTAATCCAAT 311480 CTAGAATATA CATATATTTA GAATTAACTA TTAATTATAC TAAAATATAT TATAACTAAA 3111400 CAACATTTCA AAAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAATAAAA AACCCTTAGT 311520 AGATGCAGAA AAAAAATAACC TCTTAACTCT AATAAAATAA						310860
TTTTGCANT GCTTGGGAAA CGTAAGCGCT CAAGCTCAA ACACAATAAT 311100 AAAAAAAAGA AAAAAAAAGA AAGAACCCCC GTAACATGAA AAGAGCCCAA 311100 ATAGGAGATC ATAAAATTAT CAAAAAAAGT TACCATTGA AATAAAAGCC ATTCAAAAGC 311160 TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTG TCATAATTTT TAAAATTATT TAAAATTAT 311280 CTAGAATATA CATATAACTA GTTTATTATC TTAACTTATAT TATAATTATA 311400 CTAGAATATA CATATAATTTA GAATTAACTA TTAATTATATA TATAAATTATA 311400 CAACATTCA AAAAAAAATTATTATG GCTAAAGATT AAAAAAAAAAAA 311400 CAACAATTCA AAAAAAAAAAAACC TCTTAACCTA AATAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA						310920
AAAAAAAAAA AAAGTTACTC TATTTGCCAA AGAAACTCCC GTAACATGAA AAGAGCCCAA ATAGGAGATC ATAAAAATTAT CAAAAAAAGT TACCATTTGA AATAAAAGCG ATTCAAAAGC TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTGG TCATAATTTT TAAACTTTTT 311220 GATCATATCC CTTATATACA GTCTCCTTTA TTTATTACTT TTAACTAAAT TTAAACTAAT 311280 CTAGAATATC CATATATTTA GAATTAACTA TTAAATTACC TAAAATATAT TATAATTAAA 311340 CTAGAATATA CATATATTTA GAATTAACTA TTAAATTATC TAAAATATAT TATAATTAAA 311400 CAACATTTCA AAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAGAA AAAAATAACC TCTTAACTCT AATAAATAAA AAAATACCAA TTCCTGAAAA 311580 TTATAAAAATA CCTGATCTGG TAAATATTGA TGATTTGAA GATCTAAAAA ATCTTGGACC 311640 AAAAAAAATTT GAGATCAGAA AAATATTAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAAATTT GAGATCAAAA TCAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATAATAGATGA 311940 TTACGTTTCA AACACAAAAG AAGCAATAGAAC GGACCTGGA TATAAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAAAC ATATGGAAAA 312120 ATATGCAAAC TAAATACTTC TCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGATAA 312180 ATATGCAAAC TAAATACTTC TCCGGCATT CAAATAAGAA CAAAAAAAAT TTCATAAAAC GCCCTTCCCG 312240 AACCAAGACT ATGATTCCT AAGCAGAATT ACAAAAAAAAAA						310980
TGTAGGAGATC ATAAAATTAT CAAAAAAAGT TACCATTIGA AATAAAAGCG ATTCAAAAGC TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTGG TCATAATTTT TAAACTTTTT 311220 GATCATATTC CTTATATACA GTCTCCTTTA TTTATTACTT TTAACTAAAT TTAAACTAAT 311280 CTAGAATATA CATATATTTA GAATTAACTA TAAATTATC TAAAATATAT TATAATTAAA 311340 TAATATGAAA TCAATTTATG CTTTATTATT TCTATTTATT AATTTATCTT TGTTGGCTAA 311460 CAACATTTCA AAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGAAA 311580 TTATAAAAATA CCTGATCTGG TAAATATGA TGATTTGAA GATCTTAAAA ATCTTGGACC 311640 AAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 AAAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 AATTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTAAAA ATATAGATGA 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAAC TAATATTACA 312100 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAAAT TTCTTAGATA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAAAAT TTCCTAGATAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAAAAT TTCATAAAACA GCCCTTCCCG 312240 AACTAAAGACT ATGATTCCT AAGCAGAAAT ACAAAAAAAAAA						
TGTAGGAATA GCTATAACAA ATAGCTCTTT TATTATCTG TCATAATTT TAAACTTTT 311220 GATCATATTC CTTATATACA GTCTCCTTTA TTTATTACTT TTAACTAAAT TTAATCAAAT TTAAACTAATT 311240 CTAGAATATA CATATATTTA GAATTAACTA TTAATATACT TAAAATATAT TATAAATAAA 311400 CAACATTTCA AAAAAAGATT TCTTATTATT ACTATTATATT TGTATATATT AATTAATATAA 311400 CAACATTTCA AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA						311100
CATCATATTC CTTATATACA GTCTCCTTTA TTTATTACTT TTAACTAAAT TTAATCCAAT 311280 CTAGAATATA CATATATTA GAATTAACTA TTAATTATAC TAAAATATAT TATAATTAAA 311340 TAATATGAAA TCAATTTATG CTTTATTATT TCTATTTATT TAATTTATCTT TGTTGGCTAA 311400 CAACATTTCA AAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAGAA AAAAAATAACC TCTTAACTCT AATAAATAAA AAAATCCAA TTCCTGAAAA 311580 TTATAAAAATA CCTGATCTGG TAAATATTGA TGATTTTGAA GATCTTAAAA ATCTGGAGC 311640 AAAGACTATT AAAGTAAGAA AAATATTGA TGATTTTGAA GATCTTAAAA ATCTGGAGC 311700 AAAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TTACTTTATA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTC TTCCGGCATT CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTC TTCCGGCATT CAAAAAAAAAA						311160
CTAGAATATA CATATATTTA GAATTAACTA TTAATTATAC TAAAATATAT TATAATTAAA 311340 TAATATGAAA TCAATTTATG CTTTATTATT TCTATTTATT AATTTACTT TGTTGGCTAA 311400 CAACATTCA AAAAAAGATT TAGAAGTACT GCCAAAGATT GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAGAA AAAAATAACC TCTTAACTCT AATAAATAAA AAAATACCAA TTCCTGAAAA 311580 TTATAAAAATA CCTGATCTGG TAAATATTGA TGATTTGAA GATCTTAAAA ATCTTGGAGC 311640 AAAGACTATT AAAGTAAGAA AAATATTAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAAT GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAAATTACAA 312120 ATATGCAAAC TAAATACTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAAT AAAACAAAAT CTTATTAACC TAAATTGAAAA 312120 ATCAATATCA AGCTTATCGT AAGCAGAAT AAAACAAAAT CTTATTAACC TAAATTGAAAA 312120 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAAATCAA GCTCTTCCG 312240 AACTAAAGACT ATGATTTCC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAAATATCC 312360 TGCTTGTTCA AGTCCAAAAT AATTACCAG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360	TGTAGGAATA GCTATAACA	A ATAGCTCTT	r tattatctgg	ТСАТААТТТ	TAAACTTTTT	311220
TAATATGAAA TCAATTTATG CTTTATTATT TCTATTTATT AATTTATCTT TGTTGGCTAA 311400 CAACATTCA AAAAAAGATT TAGAAGTACT GCTAAAGATT GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAGAA AAAAATAACC TCTTAACTCT AATAAATAAA AAAATACCAA TTCCTGAAAA 311580 TTATAAAAATA CCTGATCTGG TAAATATTGA TGATTTTGAA GATCTTAAAA ATCTTGGACC 311640 AAAGACTATT AAAGTAAGAA AAATATTAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311880 TAATTTACTA AACACAAAAG AAGGAAAAT GCTTTATGAA AACTCTAAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAAGACT ATGATTCCT TATCATCAAC AAAATTAAGCT ATTTTTAATTT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCTT CTTCTGCTAT 312360 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCTT CTTCTGCTAT 312360	GATCATATTC CTTATATAC	A GTCTCCTTT	A TTTATTACTT	ТТААСТАААТ	TTAATCCAAT	311280
CAACATTICA AAAAAAGATI TAGAAGTACT GCTAAAGATI GCCCAAGCAA TGAATAAGGA 311460 ATGCAAAAAAT TITATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAGAA AAAAATAACC TCTTAACTCT AATAAATAAA AAAATACCAA TTCCTGAAAA 311580 TTATAAAATA CCTGATCTGG TAAATATTGA TGATTTTGAA GATCTTAAAA ATCTTGGAGC 311640 AAAGACTATT AAAGTAAGAA AAATATAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAATTT GGGATTGAAA TTAAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCCG 312240 AACTAAAGACT ATGATTCCT TATCATCAAC AAAATTAGCT ATTTTAATT TCAAATTACC 312300 TGCTTGTTCA AGCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360	СТАGААТАТА САТАТАТТ	A GAATTAACTA	A TTAATTATAC	ТААААТАТАТ	AAATTAATAT	311340
ATGCAAAAAT TTTATTGAAA AAAATCCTAT TCAGTTCTTA AAAGAAATAA AACCCTTAGT 311520 AGATGCAGAA AAAAATAACC TCTTAACTCT AATAAATAAA AAAATACCAA TTCCTGAAAA 311580 TTATAAAATA CCTGATCTGG TAAATATTGA TGATTTTGAA GATCTTAAAA ATCTTGGAGC 311640 AAAGACTATT AAAGTAAGAA AAAATATAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGCACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312160 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTC TCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTCCT TATCACACA AAAATAAGCT ATTTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGCCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCTT CTTCTGCTAT 312360	ТААТАТСААА ТСААТТТАТ	G CTTTATTAT	TCTATTTATT	AATTTATCTT	TGTTGGCTAA	311400
AGATECAGAA AAAAATAACC TCTTAACTCT AATAAATAAA AAAATACCAA TTCCTGAAAA 311580 TTATAAAAATA CCTGATCTGG TAAATATTGA TGATTTTGAA GATCTTAAAA ATCTTGGAGC 311640 AAAGACTATT AAAGTAAGAA AAATATTAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312160 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGCCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCT CTTCTGCTAT 312360 TGCTTGTTCA AGCCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCT CTTCTGCTAT 312360	СААСАТТТСА ААААААСАТ	T TAGAAGTACT	GCTAAAGATT	GCCCAAGCAA	TGAATAAGGA	311460
TTATAAAATA CCTGATCTGG TAAATATTGA TGATTTTGAA GATCTTAAAA ATCTTGGAGC 311640 AAAGACTATT AAAGTAAGAA AAATATTAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311940 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312000 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TGCTTGTTCA AGTCCAAATT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	ATGCAAAAAT TTTATTGAA	А ААААТССТАТ	TCAGTTCTTA	AAAGAAATAA	AACCCTTAGT	311520
AAAGACTATT AAAGTAAGAA AAATATTAAT CGAAGATTTA ATTCGACTAA TAAAAGATGC 311700 AAAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCTT CTTCTGCTAT 312360	AGATGCAGAA AAAAATAAC	C TCTTAACTCT	ААТАААТАА	AAAATACCAA	TTCCTGAAAA	311580
AAAAAAATTT GGGATTGAAA TTAAAATCAA ATCTGCTTAC AGAACGCAAG AATATCAAAA 311760 ATTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AAATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360	TTATAAAATA CCTGATCTG	G TAAATATTGA	TGATTTTGAA	GATCTTAAAA	ATCTTGGAGC	311640
ATTTTATTT GATTACAATG TCAAAACTTA TGGCAGAAAA GTTGCAGAAA CCCAATCAGC 311820 AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312240 AACTAAGACT ATGATTTCC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAAATCTT CTTCTGCTAT 312360 TTTTAAAATCCA TCCAAATTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	AAAGACTATT AAAGTAAGA	ТААТТАТТААА А	CGAAGATTTA	ATTCGACTAA	TAAAAGATGC	311700
AATTCCAGGC CATTCTCAAC ATCATATGGG AACAGCAATA GATTTTATAA ATATAGATGA 311880 TAATTTACTA AACACAAAAG AAGGAAAATG GCTTTATGAA AACTCCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTTAAATCCA TCCAAATTTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	AAAAAATTT GGGATTGAA	А ТТААААТСАА	ATCTGCTTAC	AGAACGCAAG	AATATCAAAA	311760
TAATTTACTA AACACAAAG AAGGAAAAT GCTTTATGAA AACTCTCTAA AATACGGATT 311940 TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTAAATT TCAAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTTAAATCCA TCCAAATTTT CTTTTATAGT TTTTAATTGCCAG CACTTGTCAA 312420	ATTTTTATTT GATTACAATO	G ТСААААСТТА	TGGCAGAAAA	GTTGCAGAAA	CCCAATCAGC	311820
TTCCGTTTCA TACCCAAAAG GATATGAAAC GGACACTGGA TATAAAGCAG AGCCTTGGCA 312000 CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTAAAATCCA TCCAAATTTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	AATTCCAGGC CATTCTCAAG	ATCATATGGG	AACAGCAATA	GATTTTATAA	ATATAGATGA	311880
CTACTTATAC ATAGGACCTA AGCCATGCTT TATTCAGAAA AAATATTTTA ATAATTTACA 312060 ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTCTC TATCATCAAC AAAATTAGCT ATTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTAAATCCA TCCAAATTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	TAATTTACTA AACACAAAA	AAGGAAAATG	GCTTTATGAA	AACTCTCTAA	AATACGGATT	311940
ACATAAGCTT CTTGAATTTT GGAACCAGAA CAAAACAAAT CTTATTAACC TAATTGAAAA 312120 ATATGCAAAC TAAATACTTC TTCCGGCATT CAAATAAGAA CAAAAAAGAAT TGTCTAGTAA 312180 ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTCTC TATCATCAAC AAAATTAGCT ATTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTAAAATCCA TCCAAATTTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	TTCCGTTTCA TACCCAAAAC	GATATGAAAC	GGACACTGGA	TATAAAGCAG	AGCCTTGGCA	312000
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ATCAATATCA AGCTTATCGT AAGCAGAATT ATCATAAAAA AATAAATCAA GCTCTTCTCG 312240 AACTAAGACT ATGATTTCTC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTAAAATCCA TCCAAATTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	ACATAAGCTT CTTGAATTTT	GGAACCAGAA	СААААСАААТ	CTTATTAACC	TAATTGAAAA	312120
AACTAAGACT ATGATTCTC TATCATCAAC AAAATTAGCT ATTTTTAATT TCAAATATCC 312300 TGCTTGTTCA AGTCCAAATA AATTACCAGG CCCCCTTAGC CTTAAATCTT CTTCTGCTAT 312360 TTTAAAATCCA TCCAAATTTT CTTTTATAGT TTTTAATCGA AATTTGCCAG CACTTGTCAA 312420	ATATGCAAAC TAAATACTTC	TTCCGGCATT	CAAATAAGAA	CAAAAAAGAT	TGTCTAGTAA	312180
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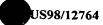
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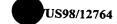


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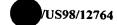
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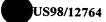
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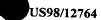


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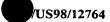
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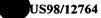


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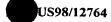


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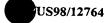


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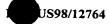
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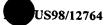
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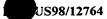
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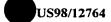


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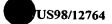
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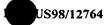
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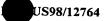
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TATTTTTCAA TAAAGCACCT CTAAAAAATG CACCAAATTA GAATAACTTT TACTAAACAT	376560
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			371			
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AGCARTTARA TATCACCCAG ACAGAARTCA AGGGARTGAA GAAGCCGCCT CTATCTTTAA 381660 AGAAGCCACT CAGGCTTACG AAATTTTAAT AGATGACAAT AAAAAAGCTA AATACGACAG 381720 ATTTGGGCAT TCCGCTTTTG AAGGAGGAG ATTTGAAGGA TTTTCAGGTG GATTTACTGG 381780 ATTTTCAGAC ATCTTGAAG ATTTTGGCGA TATTTTGAT TCATTTTTCA CTGGAAACAA 381840 AGGACAAGAA AGAAATAGAA AACACCCAAA AGGTGAAGAC TTAGGATACA ACATAGAAAT 381900 ATCTCTTGAA AATGCCATAC TTTGGGTACA AAAAATAACA TAAACATAAC AAGACAAATG 382020 ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA 382080 TGTTCTAAAAT GTTACGGAGA AGGTAAAATA ATATCAAACC CTTGTAAATC CTGTAAAGGA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCACCCCAGG CATTGATATA 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATTTAAACA TTCACACCAG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATTTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAAATTT GATAAAACTT CATAAAGTAT TCAAAAGAAA TGGTAAAGGA 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAGAA TGGTAAAGAT 382280 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382280 ATATTAAATCA CCAAAATAAA AACACCTAAA AATTCCAA AAGGAATAAA CAATGAAGAA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTCCAAA AATTACCATA TAAAACTTTTT 382660 GAAAACTTGG GCAAAAGAAT AAAAAATGA AATTCCAAA AATTACAAAG GATTATATA 382600 GTATATAACA CAAAAAAAA AACACCTAAA AATTTCAAAAAAA AATAACAAAG GATTTGAACT 382600 GTATATAACA CAAAGAATT AAAAAATGA AAGAAATGA AATTCAAAAAAAAAA	AAGAGGCTGA TTATGAGGTT GTTGACGAGG ATAAAAAATA GTGAAAAAAG ATTATTATGA	381540
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ATTTGGGCAT TCCGCTTTG AAGGAGGAG ATTTGAAGGA TTTTCAGGTG GATTTAGTGG 381780 ATTTTCAGAC ATCTTTGAAG ATTTTGGCGA TATTTTTGAT TCATTTTTCA CTGGAAACAA 381840 AGGACAAGAA AGAAATAGAA AACACGCAAA AGGTGAAGAC TTAGGATACA ACATAGAAAT 381900 ATCTCTTGAA AATGCCATAC TTTGGGTACA AAAAATAACA TAAACATAAC AAGACAAATG 381960 CTCTGTGATT CTTGTCTCGG GAAAAAATCC GAAAAAGGTA CAAGTCCTTC GATATGTAAC 382020 ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GCCGGAGGAT TTTTCAGAGT TACAACAACA 382080 TGTTCTAAAT GTTACGAGGA AGGTAAAATA ATATCAAACC CTTGTAAATC CTGTAAAGGA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACC TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTAG TAAAAATATT GATAAGATCT CATAAAGTAT TCCACAGAG AGTGAAAATA 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGGGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTCCAAA AAGGAATATA TGAAAAATTT 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTG GCAAAAGATT CAATAATTGA AAGCATAAA AATTTAAATT CTAATGCTAT TAAACATTTT 382620 GTATATAACC CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTTT 382660 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATTAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAACT CAAAACTA AAGAAATGA ACACCAAAA 382740 ATATTAAAAAT AATAAGAAAT CAAAACTA AAGAAATGA AATTCTCAAA AGCATAAATAT 382660 AAGATTTTGA AAAAAAAAAAT CAAACTA AAGAAATTAA AACACAAAA AATTCTTAAAA AACAAAACA AATAACAAAA AATTCTAAAAA AATAACAAAAG GATTTTAACTC AAAAAAAAAA	AGCAATTAAA TATCACCCAG ACAGAAATCA AGGGAATGAA GAAGCCGCCT CTATCTTTAA	381660
ATTTTCAGAC ATCTTTGAAG ATTTTGGCGA TATTTTGAT TCATTTTCA CTGGAAACAA AGGACAAGAA AGAAATAGAA AACACGCAAA AGGTGAAGAC TTAGGATACA ACATAGAAAT AGGACAAGAA AGAAATAGAA AACACGCAAA AGGTGAAGAC TTAGGATACA ACATAGAAAT ATCTCTTGAA AATGCCATAC TTTGGGTACA AAAAATAACA TAAACATAAC AAGACAAATG CTCTGTGATT CTTGTCTCGG GAAAAAATCC GAAAAAGGTA CAAGTCCTTC GATATGTAAC ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA ATGTGTAAAG GTTACCAAGACA AGGACACATT CAATTAAACA TTCCCCCAGG CATTGATAAGA AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT AACCAACAAA TAAAAATATT GATAAGATCT CATAAAGGAT TCAAAAGAAA TGGTAAAGAT CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAAGCC TTGGAAAAGA AGTGAAAATA AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA ATATTAAAC CCAAAATAAA AACACCTAAA AATTTCAAAC ACCGAAAAGTT TGGAAATTTA 382500 ATATTAAATCA CCAAAATAAA AACACCTAAA AATTTCAAAA CACAAAGGT TAAACCTTTTT 382660 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAAATAG ATTTACTCAA AGCATAATAT 382660 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT AATAAAAAAAAAACT CAAAAACTA AAGCATAAAA AATAACAAAG GATTTGAACC AATATTAAAAAAAAAAAACT CAAAAACTA AAGAAATTGA ACAGCTAACA AATTCTTAAA AACAAAAGACC AAGAGGATT TGCATTAAAA TAAAAAAATG AACTTGACAA AATTCTTAAAA AACAAAACCA 382800 ACAGAGGATT TGCATTAAAA TAAAAAATG AACTTGACAA AATTCTTAAA AACAAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAAA ACACAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTATAA AAAAAAAAAA	AGAAGCCACT CAGGCTTACG AAATTTTAAT AGATGACAAT AAAAAAGCTA AATACGACAG	381720
AGGACAAGAA AGAAATAGAA AACACGCAAA AGGTGAAGAC TTAGGATACA ACATAGAAAT 381960 ATCTCTTGAA AATGCCATAC TTTGGGTACA AAAAATAACA TAAACATAAC AAGACAAATG 381960 CTCTGTGATT CTTGTCTCGG GAAAAAACC GAAAAAGGTA CAAGTCCTTC GATATGTAAC 382020 ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA 382080 TGTTCTAAAT GTTACGGAGA AGAACCATT CAATTAAACC CTTGTAAATC CTGTAAAGGA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAATATT GATAAGATCT CATAAAGTAT TCAAAAGAAA TGGTAAAGAT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGGCC TTGGAAAAGA AGGAAAATAA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTAAA TTAAAAATGC AGGGATGCCA ATTCTCAAA AAGGAATAAA CAATGAAGAA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAA AATTACAAAG GATTTGAACT 382620 GTATATAACT CATGCCAAT CAATAATTGA AAGCATAAAA AATTACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAAC CCAAAAACTA AAGCATAAAAA AATTACTTAAA AACAAAAGACC 382800 ACAGAGGATT TGCATTAAAA TAAACAAATG AACTTGACAA AATTCTTAAA AACAAAAGACC 382800 ACAGAGGATT TGCATTAAAA TAAAACATTG AACAAAATAA AAATCAAAAA AACACAAAA 382920 ACAGAGGATT TGCATTAAAA TAAAACATTA AAAAAAAATAA AAATCTTAAAA ACCACAAAA CCA 382920 TAGATGAAAT AATAAGAATA AATACAAATG AACTTTTA AAAACAAAC AAAACCAAA 382920 TAGATGAAAT AACACTACTA GAAACATTA AAAAAAAATAA AAATCTTAAAA ACCACAAACCA 382920 TAGATGAAAT TGCATTAAAA TAAAACATTA AAAAAAAAAA	ATTTGGGCAT TCCGCTTTTG AAGGAGGAGG ATTTGAAGGA TTTTCAGGTG GATTTAGTGG	381780
ATCTCTTGAA AATGCCATAC TTTGGGTACA AAAAATAACA TAAACATAAC AAGACAAATG 381960 CTCTGTGATT CTTGTCTCGG GAAAAAATC GAAAAAGGTA CAAGTCCTTC GATATGTAAC 382020 ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA 382080 TGTTCTAAAT GTTACGGAGA AGGTAAAATA ATATCAAACC CTTGTAAATC CTGTAAAGGA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAATATT GATAAGATC CATAAAGTAT TCAAAAGAAA TGGTAAAGAT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTCCAA AAGGAATAAA CAATGAAGAA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAA AATAACAAAG GATTTGAACT 382680 TTATATATCA CAAAAAAAAAA AACACCTAAA AAGAAATGA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAAT AAAAAACTA AAGAAATTGA ACAGCTAAAA AATTCTAAAA AACACAAAA 382740 ATATAAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACACAAA 382740 ATATAAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACACAAAC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAAAAA	ATTTTCAGAC ATCTTTGAAG ATTTTGGCGA TATTTTTGAT TCATTTTTCA CTGGAAACAA	381840
ATGTGTATC CTGGTCTCG GAAAAAATCC GAAAAAGGTA CAAGTCCTTC GATATGTAAC ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA AGCAACAACA GGCAGCGGAGA AGGTAAAATA ATATCAAACC CTTGTAAATC CTGTAAAGGA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAATATT GATAAGATCT CATAAAGTAT TCAAAAGAAA TGGTAAAGAT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCCC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGG GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA 382560 GAAACCTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATTACCAAA GGATTAGAACT 382680 TTATATATCA CAAAAAAAAAAAGT CCAAAAACTA AAGAAATTGA ACACTAAAA AATTACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAAGT CCAAAAACTA AAGAAATTGA ACACTAAAA AATTCTAAA AACACAAA 382740 ATATAAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACACAAAA 382740 ATATAAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACACAAAA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTTAAAA ACCAAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTTAAAA ACCACAAACCA 382920 TAGATGAAAT AAAAAAAAAC TACAAACTTTG GAGCAAATCCT TAGAACAGCA GAACAGTTTA 382920 GTATAGATCT TGCATTACTA GAAACATTTA AAAAAAAAAA	AGGACAAGAA AGAAATAGAA AACACGCAAA AGGTGAAGAC TTAGGATACA ACATAGAAAT	381900
TGTTCTAAAT GTTACGGGAA AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA TGGTAAAGAT 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTAA TTAAAAATGC AGGGATGCCA ATTCTCAAA ACGAATATA CAATGAAGAA 382500 ATATTAAACA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382500 GAAAACTTGG GCAAAGAATT AAAAGATGG GATGAAAATA AATAACAAAG GATTTGAACT 382600 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATTACCAAAACAAAG GATTTGAACT 382600 GTATATAACA AAAAAAAAGT CCAAAAACTA AAGCATAAAA AATACCAAGA GATTTGAACT 382600 ACAGAGGATT TGCATTAAAA TATAAACTG AAAAAAATAA AAAAAAACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGCTTTC ATTTTACTTC 382920 TAGATGAAAT AGAAGATCC CAAAACTTTA AAAAAAAAAA	ATCTCTTGAA AATGCCATAC TTTGGGTACA AAAAATAACA TAAACATAAC AAGACAAATG	381960
TGTTCTAAAT GTTACGGAGA AGGTAAAATA ATATCAAACC CTTGTAAATC CTGTAAAGGA 382140 AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAATATT GATAAGATC CATAAAGTAT TCAAAAGAAA TGGTAAAGAT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 ATATTAAAAAA AATAAAAAAGT CCAAAAACTA AAGAAATTGA ACAGCTAGCT AAAACACAAA 362740 ATATAAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTT AAAAAAATAA AAATGTAAAA ACACAAAACCA 382920 AAGATTTTGA AAACTTACTA GAAACATTTA AAAAAAATAA AAATGCTAAAA ACACAAACCA 382920 TAGATGAAAT AGAAGATCC CAAAACTTTA GAGCAATCCT TAGAACAGCA GAACAGTTTAC 382920 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATTCA ACAGTTTTGC 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAATGC AGGTAATAT AAAGCAAAC 383100 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAATGC AGTGACAAAC ATAAACAAA 383100 CAATAAAATCT TTTAAAAAC TATGGCTTTT GGATATATAC TGGTGATATT AAAGGACAAG 383160 CAATAAAATC TTTAAAAAC TATGGCTTTT GGATATATAC TGGTGATATT AAAGGACAAG 3831100	CTCTGTGATT CTTGTCTCGG GAAAAAATCC GAAAAAGGTA CAAGTCCTTC GATATGTAAC	382020
AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT 382200 AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382260 GATCTTTATG TAAAAATATT GATAAGATCT CATAAAGTAT TCAAAAGAAA TGGTAAAGAT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382680 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAGT CCAAAAACTA AAGCATAAAA AATTCTTAAA AACAAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAAA ACACAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAAA ACACAAACCA 382920 TAGATGAAAT AGAAGATCC CAAAACTTA AAAAAAAAGA AAATGTAAAAA ACACAAACCA 382920 TAGATGAAAT AGAAGATCC CAAAACTTTA AAAAAAAAGA AAATGTAAAAA ACACAAACCA 382980 GTATAGATGAAT AGAAGATCC CAAAACTTTG GAGCAATCCT TAGAACAGCA GAACAGTTTA 382980 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATCCA ACAGTTTTGC 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAATGA AGGACAAAC ATAAACAACA AAGATTTTGA AAACTTACT ACTCAAAAAC GCAGCGCAAA AGACAATCCA ACAGTTTTGC 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAATGA AGGACAAAC ATAAACAACA 383100 CAATAAAATCT TTTAAAAAC TATGGCTTTT GGATATTAC TGGTGATATT AAAGGACAAG 3831160 CAATAAAATCT TTTAAAAAATA AACCGATAAAA AAAATTAC TGGTGATATT AAAGGACAAG 3831160	ATGTGTAACG GCAGCGGAAG AGTAGTGCAA GGCGGAGGAT TTTTCAGAGT TACAACAACA	382080
AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT 382320 GATCTTTATG TAAAAATATT GATAAGATCT CATAAAGTAT TCAAAAGAAA TGGTAAAGAT 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAAATTTA 382560 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 ATATAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAACCA 382920 TAGATGAAAT AGAAGATCC CAAAACTTA AAAAAAAAGA AAATGCTTTC ATTTTACTTC 382920 TAGATGAAAT AGAAGATCC CAAAACTTT GAGCACAAACCA GAACCATTTC ATTTTACTTC 382980 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATTCA ACAGTTTTG 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAAATGA CAGTGACAAAC ATAAACAACA 383100 CAATAAAACCT TTTAAAAAAC TATGGCTTTT GGATATATAC TGGTGATATT AAAGGACAAG 383160 CAATAAAACCA AATCAAAATA AACGATAAAA AAATTGCACT TATTTTAGGA AATGAAGGAA 3831200	TGTTCTAAAT GTTACGGAGA AGGTAAAATA ATATCAAACC CTTGTAAATC CTGTAAAGGA	382140
GATCTTTATG TAAAAATATI GATAAGATCI CATAAAGTAI TCAAAAGAAA TGGTAAAGAI 382320 CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTAA TTAAAAATGC AGGGATGCCA ATTCTCAAA CCGAAAAGTT TGGAAATTTA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTAAAATT CTAATGCTAI TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGG GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 ATATAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAACCA 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGCTAAA ACCAAAACCA 382920 TAGATGAAAT AGAAGATCC CAAAAACTTA AAAAAAAAGA AAATGCTTTC ATTTTACTTC 382920 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATCA ACAGTTTTA 383040 GCACTAGCTC TGGCGCAAG CAATATGTAA AAAAAAATGC AGTGACAAAC ATAAACAACA 383100 CAATAAAACC TTTAAAAAAC TATGGCTTTT GGATATATAC TGGTGATATT AAAGGACAAG 383160 CAATAAAACCA AATCAAAAAA AAATTGCACT TATTTTAGGA AATGAAGGAA 3831220	AAAGGAAGTC TTACAAAGCA AGAAACCATT CAATTAAACA TTCCCCCAGG CATTGATAAT	382200
CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA 382380 AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACC CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAGT CCAAAAACTA AAGAAATTGA ACAGCTAGCT AAAACACAAA 382740 ACAGAGGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AATTCTTAAA ACCAAAACCA 382800 ACAGAGGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAAA ACACAAACCA 382860 AAGATTTTGA AAACTTACTA GAAACATTTA AAAAAAAAAA	AACCAACAAA TAAAAATGAA AGGCAAGGGA AATGTTAATC CAGATAATCA AGAATATGGT	382260
AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA 382440 CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAGT CCAAAAACTA AAGAAATTGA ACAGCTAGCT AAAACACAAA 382740 ATATAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAAATAA AAATGTAAAA ACAAAACCA 382860 AAGATTTTGA AAACTTACTA GAAACATTTA AAAAAAAAGA AAATGCTTTC ATTTTACTTC 382920 TAGATGAAAT AGAAGATCC CAAAACTTTG GAGCAATCCT TAGAACAGCA GAACAGTTTA 382980 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATTCA ACAGTTTTGC 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAAATGAC AGTGACAAAC ATAAACAACA 383100 CAATAAAACT TTTAAAAAAC TATGGCTTTT GGATATTAC TGGTGATATT AAAAGGACAAG 383160 ATATAAAACAA AATCAAAATA AACGATAAAA AAATTGCACT TATTTTAGGA AATGAAGGAA 383120	GATCTTTATG TAAAAATATT GATAAGATCT CATAAAGTAT TCAAAAGAAA TGGTAAAGAT	382320
CAAAATTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA 382500 ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAGT CCAAAAACTA AAGAAATTGA ACAGCTAGCT AAAACACAAA 382740 ATATAAAAAA AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGGATT TGCATTAAAA TTAAAACTTG AAAAAAAATAA AAATGTTAAAA ACAAAACCA 382860 AAGATTTTGA AAACTTACTA GAAACATTTA AAAAAAAAAA	CTCTATGCAA TGCTTCCAAT AAGCTTTACT CAAGCAGCGC TTGGAAAAGA AGTGAAAATA	382380
ATATTAATCA CCAAAATAAA AACACCTAAA AATTTAAATT CTAATGCTAT TAAACTTTTT 382560 GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAGT CCAAAAACTA AAGAAATTGA ACAGCTAGCT AAAACACAAA 382740 ATATAAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAAATAA AAATGTAAAAA ACACAAACCA 382860 AAGATTTTGA AAACTTACTA GAAACATTTA AAAAAAAAGA AAATGCTTTC ATTTTACTTC 382920 TAGAATGAAAT AGAAGATCC CAAAACTTTG GAGCAAATCCT TAGAACAGCA GAACAGTTTA 382980 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATTCA ACAGTTTTGC 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAAATGAC AGTGACAAAC ATAAACAACA 383100 CAATAAAATCT TTTAAAAAAC TATGGCTTTT GGATATATAC TGGTGATATT AAAGGACAAG 383160 ATATAAACAA AATCAAAATA AACGATAAAA AAATTGCACT TATTTTAGGA AATGAAGGAA 383220	AAAACAATCG CTTCAAAAGA GATTAAAATA CACATTCCAA AAGGAATAAA CAATGAAGAA	382440
GAAAACTTGG GCAAAGAATT AAAAGATGGT GATGAAATAG ATTTACTCAA AGCATAATAT 382620 GTATATAACT CATGCCAATT CAATAATTGA AAGCATAAAA AATAACAAAG GATTTGAACT 382680 TTATATATCA AAAAAAAAGT CCAAAAACTA AAGAAATTGA ACAGCTAGCT AAAACACAAA 382740 ATATAAAAAT AATAAGAATA AATACAAATG AACTTGACAA AATTCTTAAA AACAAAGACC 382800 ACAGAGGATT TGCATTAAAA TTAAAACTTG AAAAAAATAA AAATGTAAAA ACACAAACCA 382860 AAGATTTTGA AAACTTACTA GAAACATTTA AAAAAAAAAGA AAATGCTTTC ATTTTACTTC 382920 TAGATGAAAT AGAAGATCCC CAAAACTTTG GAGCAATCCT TAGAACAGCA GAACAGTTTA 382980 GTATAGATCT TGTAATTACT ACTCAAAAAC GCAGCGCAAA AGACAATTCA ACAGTTTTGC 383040 GCACTAGCTC TGGCGCAAGT CAATATGTAA AAAAAAATGAC AGTGACAAAC ATAAACAACA 383100 CAATAAAATCT TTTAAAAAAC TATGGCTTTT GGATATATAC TGGTGATATT AAAGGACAAG 383160 ATATAAACAA AATCAAAATA AACGATAAAA AAATTGCACT TATTTTAGGA AATGAAGGAA 383220	CAAATTTTAA TTAAAAATGC AGGGATGCCA ATTCTTCAAA CCGAAAAGTT TGGAAATTTA	382500
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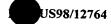
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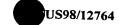
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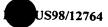
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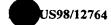


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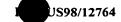


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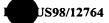
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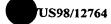
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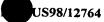
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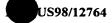


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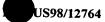
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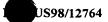
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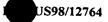
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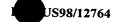


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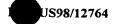
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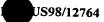


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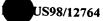


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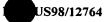
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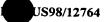
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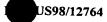
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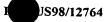
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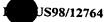
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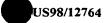
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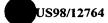
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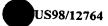


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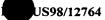
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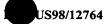


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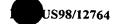


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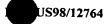
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GCTTGAGTAT AGCAACGATG TTCTTTATAA TAATAAAAAT AGGGTTGGAC ATCGTCCTTT 502020 AAAATTGGTT GCCAAAAGTA TTTATGGTAA AAATAATACA GATATTATAC TTGATGAGTA 502080 TTCTATTAAT AAGTTGTTT CAAATAGTAA CAACATTAAA CTCTGCAAG ATGGAAAACT 502140 TGTTGTAATA AAGTAAGAAA ATTAAATTA GTACTTGTTT TTTCTAAAC AATCAACTAA 502200 GATAATAAAG AGTTTTATAG GGAATTAGTG CCCTTATAGC TCAGTTGGTA GAGCACCACC 502260 ATGGTAAGGT GGGGGTCTC GGTTCAAGTC CGATTGAGG CTTTTATTGG TAGTTAGGAG 502320 TTTTGTTATA TGGGTAAAAA GAAGGCAAA GGAGCTGTTG AGCTTATATC TTTGATTTGT 502380 GAAGAAACAG GAATTAGAAA TTATACCACT ACTAAGAATA GACGCAATAA GCAAGAAAAG 502440 TTAGAATTGA TGAAAATATTG TCCAAAATTA CGAAAACACA CTCTTCATAA AGAAGGAAAA 502500 ATGCTGGGGG TTCGAATCCC TCCTGACCTC TTGTTAGAAT TAAAGTGGGC CTAAGATATAAT 502620 TTTAGGTTTA TCAAAGATGG TATCTTAGGG CTTAAGAAGG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTTT TGGCTGGTAT TATTTTGTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTCTTGTT GTAACTTATG TATTTTAGTT TAATTATATAA 502880 AAGGTTAAGC GAAATAAG GCTTGTATA TATTTTAGTT TATTTTATTT	TAATACTTCT	GGTTCTAAAA	AATTAGAAG	A CTCTTTTTT	T ATCAAAATT1	r atgatgaaaa	501900
AAAATTGGTT GCCAAAAGTA TTTATGGTAA AAATAATACA GATATTATAC TTGATGAGTA 502000 TTCTATTAAT AAGTTGTTT CAAATAGTAA CAACATTAAA CTTCTGCAAG ATGGAAAACT 502140 TGTTGTAATA AAGTAGAAA ATTAAATTTA GTACTTGTTT TTTTCTAAAC AATCAACTAA 502200 GATAATAAAG AGTTTTATAG GGAATTAGTG CCCTTATAGC TCAGTTGGTA GAGCACCACC 502260 ATGGTAAGGT GGGGGTCGTC GGTTCAAGTC CGATTGAGGG CTTTTATTGG TAGTTAGGAG 502320 TTTTGTTATA TGGGTAAAAA GAAGGGCAAA GGAGCTGTTG AGCTTATATC TTTGATTTGT 502380 GAAGAAACAG GAATTAGAAA TTATACCACT ACTAAGAATA GACGCAATAA GCAAGAAAAG 502440 TTAGGAATTGA TGAAATATTG TCCAAAATTA CGAAAACCAC CTCTTCATAA AGAAGGAAAA 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTC TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502620 ATGCTGGGGG TTCGAATCCC TCCTGACCTC TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502620 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTC GAAATGGAAA GCAAGTTTTT TGGCTGGTAT TATTTGTTTC AATTTTCTTG 502740 GGTATAGTCC ATTACTTAT GTTTCTTGTT GTAACTTATG TATTTTTATATAG 502800 AAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGGTCTAAAAC TTATTCTCAA TATGAGAAAA 502820 AGGAGAAAAAT TATGTCTAGA GCTTGGTATG TAGGTCAAAC TTATTCTCAA TATGAGAAAA 502920 ATGTTAAGGC GGACATAAGG CTTTTAATAA ATGAAGGTGT TTTTGGCTGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAG AGATAAGAAA TGGCAAGAAA AGAATAAGGG 502920 ACGAGAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTGTA TCTTCCAGA GTAGGCTGGA 503040 AAGATATAT TGCTAATATT ATCAAAGTTC AAGGCTGTA TATTTTTTT TGGTTGTTT GTGTTTAGTA 503100 AAGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAGCTATA AAGGGTTAT TATTTTTTT TGGTTTAGTA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAGCTATA AAGGGTTTT TATGTTTTT TTGTTTTTT TTGGTTTAGTG 503160 GTGAGAATAA AGGAATAAA TCTATTTTTT TGGCTTTAGA TCTTTAGATCT ATTGATTTAT TGCTTACTG 503140 ATTCCAACA TATAGAAAAA TCTATTTTTT TGAAGAGAC TTATTAGTTCT ATTGATTATG 503340 ATTACAAAGAA AATAAAAAA TCTATTTTTTA TGCTTTATAG CTTTTGAACCCCT GTTGAAGTTG 503340 ATTACAATAAA GGAAATAAA ATTAAAATT TAATACTAAT GGGAGCTTTG AAGGCGTTAC 503340 ATTTCCAACA TATAGAAAAA ATTAAAAAT TAAAATTT TAATACTAAT TTGGATTAAA TTGCAGGTTC 5033400 ATTCCCAAAAAAAAA ATTAAAAAAAAAAAAAAAAAAAA	CATTAGGCCA	TATTTTGATA	AGAGAATGG	I TTCTTCAGA(G GCTTTGAAA!	AATGGGGAAT	501960
TTCTATTAAT AAGTTGTTTT CAAATAGTAA CAACATTAAA CTTCTGCAAGA AATCAACTAA 502200 GATAATAAAG AAGTAAGAAA ATTAAATTA GTACTTGTTT TTTTCTAAAC AATCAACTAA 502200 GATAATAAAG AGGTTATAGG CCCTTATAGC TCAGTTGGTA GAGCACCACC 502260 ATGGTAAGAG GGGGGTCGTC GGTTCAAGTC CGATTGAGGG CTTTTATTGG TAGGTTAAGAA 502320 TTTTGTTATA TGGGTAAAAA GAAGGCAAA GGAGCTGTTC ACCACAATAA GCAAGAAAAG 502440 TTAGAATTGA TGCAAAATTA CCAAAACACA CTCTCATAAA ACGACAGTCT CCAAAACTG 502500 ATGCTGGGG TTCGAACCC TCCTGACCTC TTGTTAGAAG CTTAAAGAGG CTTTAAAGTG 50260 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAGG TATATTTTTC 502740 GGTATAGTCG ATTATCTTAT TGGCTGGTAT TATTTTTTTA 502740 GGTATAGTCG ATTATCTTAT TGGTCTGAAA TATTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT	GCTTGAGTAT	AGCAACGATG	TTCTTTATA	A AAAAATAAT	T AGGGTTGGAC	C ATCGTCCTTT	502020
GATAATAAAG AGTTTATAG GGAATTAGT CCCTTATAGC TCAGTTGGTA GAGCACCACC 502260 ATGGTAAGGT GGGGGTCGTC GGTTCAAGTC CGATTGAGGG CTTTTATTGG TAGTTAGGAG 502320 TTTTGTTATA TGGGTAAAAA GAAGGGCAAA GGAGCTGTTG AGCTATATC TTTGATTTGT 502380 GAAGAAACAG GAATTAGAAA TTATACCACT ACTAAGAATA GACGCAATAA GCAAGAAAAG 502440 TTAGAATTGA TGAAATATTG TCCAAAAATTA CGAAAACACA CTCTTCATAAA AGAAGGAAAA 502500 ATGAAATAAT AAATAATAGG TCAGTAGTTC CAACGGTAGA ACGACAGTCT CCAAAACTGT 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTC TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502620 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAAG TAACTTCTCTTG 502740 GGAAGTTGTTG GAAATCGAA GCAAGTTTTT TGGCTGGTAT TATTTGTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTATG TATTTTAGTT TATTATAAAG 502800 AAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTGTTC AATTTTCTTG 502920 ATGTTAAGGC ACCATAAGG CTTTAATAAA ATGAAGGTGT TTTTTGGCGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAA AGAATAAGAG TCCTCAAAACTGT 502920 AGGAAAAAAT TATGCCAAGG CTTTTAATAA ATGAAGGTGT TTTTTGGCGGT GTGGTATTAG 502920 AGGAAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GAAATAAGGG 502980 AGGAAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTTGA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAGCTTAT TAATTTTTTT GGTGTTAGTA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAGCTTAT TAATTTTTTT GTGTTTAGTA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAGCTTATA CTTTCAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGCAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGCAAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGCAAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGAA GGAGAAAGAG 503320 TTAGAATTAA AGCAAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGGAC TTTTGAAGTTG 503340 ATTTCCAACA TATAGAAAAG ATTTAAAATT TAATACTAAT GGGAGCTTTG AAGGCGTTAC 503400 AATTTCCAACA TATAGAAAAAA ATTAAAAAAAA AAGCAATTC TTGGAGCCT CATGGAGTTA 503340 ATTTCCAACA TATAGAAAAAAAAAAAAAAAAAAAAAAAA	AAAATTGGTT	GCCAAAAGTA	TTTATGGTA	A AAATAATACA	A GATATTATAC	TTGATGAGTA	502080
GATAATAAAG AGTTTTATAG GGAATTAGTG CCCTTATAGC TCAGTTGGTA GAGCACCACC ATGGTAAGGT GGGGGTCGTC GGTTCAAGTC CGATTGAGGG CTTTTATTGG TAGTTAGGAG 502320 TTTTGTTATA TGGGTAAAAA GAAGGGCAAA GGAGCTGTTG AGCTTATATC TTTGATTTGT 502380 GAAGAAACAG GAATTAGAAA TTATACCACT ACTAAGAATA GCACAATAA GCAAGAAAAG 502440 TTAGAATTGA TGAAATATTG TCCAAAAATTA CGAAAACACA CTCTTCATAA AGAAGGAAAA 502500 ATAAAATAAT AAATAATAGG TCAGTAGTTC CAACGGTAGA ACGACACTCT CCAAAACTGT 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTG TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG GAAGTTGTTA TCAAAGATAG TATCTTAGAG CTTAAGAAG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTT TGGCTGGTAT TATTTTGTTTC AATTTTCTTG 502740 GGTATAGCC ATTATCTTAT GTTTCTTGTT GTAACTTATG TATTTTAGTT TTATTATAAG AGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTTTTGGCGGT GTGGTATTAG AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAGGGTT TTTTGGCGGT GTGGTATTAG AGAGAAAAAT TTGGCCAGGC TATATCTTA TTGAGCTAGA TCTCCAGAA GTAGCACGA AGAGAAAAAT TTGGCCAGGC TATATCTTA TTGAGCTAGA TCTCCAGAA GTAGCCTCGA AGAGAAAAAT TTGGCCAGGC TATATCTTA TTGAGCTAGA TCTTCCAGAA GTAGCCTCGA AGAGAAAAAT TGGCCAGGC TATATCTTA TTGAGCTAGA TCTTCCAGAA GTAGCCTCGA AGAGAAAAAT TGGCCAAGAA ACAGTACAA AGAGTAAAA AAGATTAAT GCTAATATT ACCAAAGTTC AAGAGCTTAT TAATTTTTTT TTGTTTTTTTTTT	TTCTATTAAT	AAGTTGTTTT	CAAATAGTA	A CAACATTAAA	CTTCTGCAAG	ATGGAAAACT	502140
TTTTGTTATA TGGGTAAAAA GAAGGCAAA GAAGTATTA GACGCAATAA GCAAGAAAAA GAAGGCAAA GAAGCAATAA GAAGGAAAAA GAAGGCAAA GAAGCAATAA GAAGAAAAAA TTATACCACT ACTAAGAATA GACGCAATAA GCAAGAAAAA 502500 ATAAAATAT AAATAATAG TCCAAAATTA CGAAAACAA CTCTTCATAA AGAAGGAAAA 502500 ATAAAATAAT AAATAATAG TCCAAAACTGT CAACGGTAGA ACGACAGTCT CCAAAACTGT 502560 ATAAAATAAT AAATAATAGG TCCGTGACCTG TTGTTAGAAT TAAAGTGGG CTTTAAAGTG 502620 TTTAGGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAA GCAAGTTTT TGGCTGGTAT TATTTGTTTC AATTTTCTTG 502740 GGTATAGGCG ATTATCTTAT GTTTCTTGTT GTAACTTAG TATTTTAGTT TTATTATAAG 502800 AGGGTAAAT TATGTCTAGA GCTTGATAG TAGTTCAAAC TTATTCTCAA TATGAGAAAA 502800 AGGACATAAGG CTTTAAAAG ACTTTTAGAGAAA AGAATAAGG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAA AGAATAAGAA TGGCAAGAAA AGAATAAGG 502920 AGGACATAAGG CTTTTAATAA ATGAAGGTGT TTTTGGCGGT GTGGTATTAG 503900 AAGATAATAT TTGGCCAGGC TATATCTTA TTGAGCAGAA TCTTCCAGAA GTAGGCTGGA 503040 AGAGAAAAAT TTGGCCAGGC TATATCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGAT AAGAGTAAA AAGTGTTTT TATCTTATTTGT GTGTAACTT 503100 AGGGGCAAAA AGCAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GAAGAAAAAA AGCAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GAAGAAAAAA AGCAAATAAA TCTATTTTTA TGCTTAATGA CTTTGAAGAA GAAGAAAAAA AGCAAATAAA TCTATTTTTA TGCTTAATAG CTTTGAAGAA GAAGAAAAAA AGCAAATAAA TCTATTTTTA TGCTTAATAG CTTTGAAGAA GAAGAAAAAA AGCAAATAAA TCTATTTTTA TGCTTAATAG CTTTGAAGAA GAAGAAAAAA AGCAAATAAA TCTATTTTTA TGCTTAATA CGGAGCTTTA AAGGCGTTAC 503340 AAAGAAAAAA ATTAAAAAGTT GCAGTTCAAA TTTTTGGAAGAC TATTAGTTAT ATGCAGGTT CTTGAAGAC TATTAGAAAAAAAAAA	TGTTGTAATA	AAGTAAGAAA	ATTAAATTTA	GTACTTGTT1	TTTTCTAAAC	AATCAACTAA	502200
GAAGAAAAA TATGCTAGAA GAAGGCAAA GGAGCTGTTG AGCTTATATC TTTGATTTGT 502380 GAAGAAACAG GAATTAGAAA TTATACCACT ACTAAGAATA GACGCAATAA GCAAGAAAAG 502440 TTAGAATTGA TGAAATATTG TCCAAAATTA CGAAAACACA CTCTTCATAA AGAAGGAAAA 502500 ATAAAATAAT AAATAATAGG TCAGTAGTTC CAACGGTAGA ACGACAGTCT CCAAAACTGT 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTG TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502660 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTT TGGCTGGTAT TATTTGTTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTATT TATTTAGTT TTATTATAAG 502860 AGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTCTCAA TATGAGAAAA 502860 AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAGGTGT TTTTGGCGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAA AGGATAACAAA TGGCAAGAAA AGAATAAGGG 502980 AGAGAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGAGGAAAAAT TGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGAGGAAAAAA TGGCAAGAAA ACGAATAAA TCTATTTTTA TGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAGTAAA AAGTTTTTT ATGCTTACTG 503160 GTGAGATTAA AGCAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GAGAAAAAAA ACGAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GAGAAAAAAA AAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTGGAGCT TATTAGTTCT ATTGATTATG 503280 AAAGAAAAAA ATTAAAAGTT GCAGTTCAAA TTTTTGGAGC TTATTAGTTCT ATTGATTATG 503280 ATTCCAACAA TATAGAAAAA ATTAAAATT TAAATATTATA GGGAGCCTTTG AAGGCCTTAC 503400 TACCATAAGG AGATTATAT GCAAAAAAAA AACCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520	GATAATAAAG	AGTTTTATAG	GGAATTAGTO	G CCCTTATAGO	TCAGTTGGTA	GAGCACCACC	502260
GAAGAAACAG GAATTAGAAA TTATACCACT ACTAAGAATA GACGCAATAA GCAAGAAAAG 502440 TTAGAATTGA TGAAATATTG TCCAAAATTA CGAAAACACA CTCTTCATAA AGAAGGAAAA 502500 ATAAAATAAT AAATAATAGG TCAGTAGTTC CAACGGTAGA ACGACAGTCT CCAAAACTGT 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTG TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502620 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAGG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTTT TGGCTGGTAT TATTTGTTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTAGT TATTTTAGTT TTATTATAAG 502860 AAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTCCAA TATGAGAAAA 502860 AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAGGTGT TTTTGGCGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAA AGGAATAAGAAA TGGCAAGAAA AGAATAAGGG 502980 AGAGAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGAT AAGAAGTAA AAGTGTTTTT ATGCTTACTG 503160 GTGAGATTAA AGCAAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGAA GGAGAAAGAG AAAGAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGAA GAGAAAAAAA AAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTGAAGCT TATTAGTTCT ATTGATTATG 503280 AAAGAAAAAA ATTAAAAGTT GCAGTTCAAA TTTTTGAAGCT TATTAGTTCT ATTGATTATG 503280 AAAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTGAAGGACT TATTAGTTCT ATTGATTATG 503280 ATTTCCAACA TATAGAAAAA ATTAAAATT TAATACTAAT GGGAGCTTTG AAGGCCTTAC 503400 ATTCCAACAA TATAGAAAAA ATTAAAAATA TAATACTAAT GGGAGCTTTG AAGGCCTTAC 503400 TACCATAAGG AGATTATTG GCAAAAAAAA AAGCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520	ATGGTAAGGT	GGGGGTCGTC	GGTTCAAGTC	CGATTGAGGG	CTTTTATTGG	TAGTTAGGAG	502320
TTAGAATTGA TGAAATATTG TCCAAAATTA CGAAAACACA CTCTTCATAA AGAAGGAAAA 502500 ATAAAATAAT AAATAATAGG TCAGTAGTTC CAACGGTAGA ACGACAGTCT CCAAAACTGT 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTG TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502620 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTTT TGGCTGGTAT TATTTTGTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTATG TATTTTAGTT TATTTATAAG 502800 AAAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTCTCAA TATGAGAAAA 502860 AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAGGTGT TTTTGGCGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAG AGATAAGAAA TGGCAAGAAA AGAATAAGGG 502980 AGAGAAAAAT TTGGCCAGGC TATATCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGAGGAAAAAT TTGCTAATATT ATCAAAGTTC AAGGCGTTAT TAATTTTTTT TGGTTTAGTA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAAGTAAA AAGTGTTTTT ATGCTTACTG 503160 GTGAGATTAA AGCAAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGGCGGACCT TTTGACTCCT TTGAAGGACT TATTAGTTCT ATTGATTATG 503280 AAAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTTGGAAG ATCAACGCCT GTTGAAGTTG 503340 ATTTCCAACA TATAGAAAAG ATTTAAAATT TAATACTAAT GGGAGCTTTG AAGGCGTTAC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA AAGCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA AAGCAATTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA AAGCAATTTC TTGGATTGAAC TCCGGCATTG 503520	TTTTGTTATA	TGGGTAAAAA	GAAGGGCAAA	GGAGCTGTTG	AGCTTATATC	TTTGATTTGT	502380
ATAAAATAAT AAATAATAGG TCAGTAGTTC CAACGGTAGA ACGACAGTCT CCAAAACTGT 502560 ATGCTGGGGG TTCGAATCCC TCCTGACCTG TTGTTAGAAT TAAAGTGAGG CTTTAAAGTG 502620 TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAGG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTT TGGCTGGTAT TATTTGTTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTAGT TATTTTAGTT TTATTATAAG 502800 AAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTCTCAA TATGAGAAAA 502860 AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAGGTGT TTTTGGCGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAG AGATAAGAAA TGGCAAGAAA AGAATAAGGG 502980 AGAGAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AAGGGGCAAAG GCCTATTCCT ATTAATGAT AAGGCGTTAT TAATTTTGTT GGTGTTAGTA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAAGTAAA AAGTGTTTTT ATGCTTACTG 503160 GTGAGATTAA AGGCAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGGCAGACCT TTTGACTCCT TTGAAGGACT TATTAGTTCT ATTGATTATG 503280 AAAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTGGAAG ATCAACGCCT GTTGAAGTTG 503340 ATTTCCAACA TATAGAAAAG ATTTAAAATT TAATACTAAT GGGAGCTTTG AAGGCGTTAC 503340 TACCATAAGG AGATTAATG GCAAAAAAAA AAGCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520 GTGGTCCTCA GTTGTAAAG GAATTAAAT TAAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520	GAAGAAACAG	GAATTAGAAA	TTATACCACT	' ACTAAGAATA	GACGCAATAA	GCAAGAAAAG	502440
TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTTT TGGCTGGTAT TATTTGTTTC AATTTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTAGT TATTTTAGTT TTATTATAAG 502800 AAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTCTCAA TATGAGAAAA 502860 AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAGAGT TTTTGGCGGT GTGGTATTAG 502920 ATGTTAAGGC TCCTATTGAA AAAGTAGAAG AGATAAGAAA TGGCAAGAAA AGAATAAGGG 502980 AGAGAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503040 AAGGAAAAAAT TGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAAGTAAA AAGTTTTTT ATGCTTACTG 503160 GTGAGATTAA AGGCAAATAAA TCTATTTTTA TGCTTATGA CTTTGAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGGCGACCT TTTGACTCCT TTGAAGGACT TATTAGTTCT ATTGATTATG 503280 AAAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTTGGAAG ATCAACGCCT GTTGAAGTTG 503340 ATTTCCAACA TATAGAAAG ATTTAAAATT TAATACTAAT GGGAGCTTTG AAGGCGTTAC 503400 TACCATAAGG AGATTATAT GCAAAAAAAA AAGCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520 GTGGTCCTCA GTTTGAAAG GAATTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520 GTGGTCCTCA GTTTGTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520	TTAGAATTGA	TGAAATATTG	ТССААААТТА	CGAAAACACA	CTCTTCATAA	AGAAGGAAAA	502500
TTTAGGTTTA TCAAAGATAG TATCTTAGAG CTTAAGAAGG TAACGTGGCC TAAGTATAAT 502680 GAAGTTGTTG GAAATGGAAA GCAAGTTTT TGGCTGGTAT TATTTGTTC AATTTCTTG 502740 GGTATAGTCG ATTATCTTAT GTTTCTTGTT GTAACTTATG TATTTTAGTT TTATTATAAG 502800 AAGGGTAAAT TATGTCTAGA GCTTGGTATG TAGTTCAAAC TTATTCTCAA TATGAGAAAA 502860 AGATAGAGCA GGACATAAGG CTTTTAATAA ATGAAAGATA TGGCAAGAAA AGAATAAGGG 502980 AGAGAAAAAT TTGGCCAGGC TATATTCTTA TTGAGCTAGA TCTTCCAGAA GTAGGCTGGA 503040 AAGATATTAT TGGCTAATATT ATCAAAGTTC AAGGCGTTAT TAATTTTTTT ATGCTTAGTG 503100 AGGGGCAAAG GCCTATTCCT ATTAATGATG AAGAAGTAAA AAGTTTTTT ATGCTTACTG 503160 GTGAGATTAA AGCAAATAAA TCTATTTTTA TGCTTTATGA CTTTGAAGAA GGAGAAAGAG 503220 TTAGAATTAA AGGCGGCCT TTTGACTCCT TTGAAGGACT TATTAGTTCT ATTGATTATG 503280 AAAGAAAGAA ATTAAAAGTT GCAGTTCAAA TTTTTGGAAG ATCAACGCCT GTTGAAGTTG 5033400 ATTTCCAACA TATAGAAAAG ATTTAAAAATT TAATACTAAT GGGAGCTTTG AAGGCGTTAC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAAA AAGCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAAA AAGCAATTTC TTGGATTAAA TTGCAGGTTC 503460 CAGCTGCTCA AGCGGCTCCA GGAGCTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520 GTGGTCCTCA GTTGTAAAG GAATTAAAA TAGGGCAAGC GCTTGGACCT CATGGAGTTA 503520	ATAAAATAAT	AAATAATAGG	TCAGTAGTTC	CAACGGTAGA	ACGACAGTCT	CCAAAACTGT	502560
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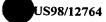


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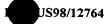
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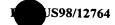
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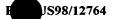
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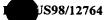
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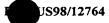
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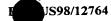
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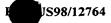
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AGTTTAAATC	CTACAGAATA	GGTCTTGTTT	TTTATAAAGA	CTATCTTGAA	GATTTTTAA	580800
CCAAAGCTTT	TGATTTTAAT	ACTATTCCTT	ATTTAAATAA	TATTCTTAAG	TATGTTAATG	580860
TTGGTGGCGG	TGGGGATTAT	CCAGAAGCTG	TTTTTGAGGG	GATTGATGCT	GCTGTGACCC	580920
AATTTGATTG	GCGGGCAGAA	AGAAGGTTTA	TTATTGTTAT	AGGAGATGCA	CCTCCTCATG	580980
AGTATCCAAG	AGGGTCTATT	GTTTATAAAG	ATGTTATCAA	TTCTGCAAAG	GAAAAAGATA	581040
TTACAATTTA	TGGAATAATA	TTTCAGTAAA	AATTTTTATT	TCTTAAATTA	TTATTTTTA	581100
TTATTTTCTA	TTTTATTTAA	TATTTTTTTA	GCTAAAGGCT	TAATCCATTT	AGGCATTTCC	581160
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TAGTATATGA	ATGCAATTTC	ATATTTACCA	GTAGCAACTA	TATTTGAATT	GTGAGCGAAA	581280
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TTTAGAATGT	TTTTATTATT	TAATTTTATT	AGATATAAAA	TAAGAAAGCA	AGGCTTTTTA	581520



AAGCCTTGCT	TTATGATTGT	TTTTGCTTAT	TTGGCAGGAA	ТТАТТАТСТТ	CCAGTTAGAA	581580
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AATTCATTGG	CAAGTCTAAA	GTTTTTTACA	TTTTTAGCCT	CAACCCCTTT	TTCCCACAAT	581820
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GCTATTTCCA	CTTTTTCTTC	GTATGCGAGT	ACTATTGGAA	TTGAAATTTC	TGCTTCTCCA	581940
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GATTTGCAAG	ACGCAAAAAT	ATATTTATTT	AGAGGAAGTG	TTGATGCTAT	AATTCATAGC	583260

			487			
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ACTTCAAAAT	TTTATGTCAA	TAGCGATTAT	TTTAATAATT	GTTTTATTAA	TGTTTACAAG	583380
GGTAGCATCA	GACATATTAA	TAAAACCGAA	TATTTAATTT	TTCCCAATAC	CAGTCTTTTG	583440
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ATGAAAGATT	ТТАААТТТАА	TTCTATATAT	TCAAAATGGA	GCCTGGAAGA	TAAAAATTAT	583680
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AAATATTTT	CTACTTTTTT	TGACAATTCA	ACAACTGTTA	ACGGGCCCAT	TTCAAAATTT	583860
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TAGGATATTA	TGTAGGTCTT	TTTTTAAAGG	AATTTATTAA	ATCTGGCTTT	GGAAATGCTG	584940
CTTTGATTGC	AGGTCAAAAT	TATCCCGTTA	TGAATGATTA	TATTTATCGT	TATTTCAAGA	585000
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1	PAATACTTCC	TATTGTGGGG	CCTGCTGTTG	AAGGAGTGCT	TTCTTCTGTT	AGAGAGAATA	585180
F	TATCTCTGC	AGTTCTTTTT	GATAGCGAAG	ATTATTTGGA	TAATAAAGAA	AATATTATTG	585240
C	STTCAGGAAT	ТАСАААТСАА	AAATATTATG	TTTCACATAT	TTTAGATAAG	GCTCTTAAGT	585300
C	CAGAGATTAA	CTATGGAAAT	TCTGATATTT	TTGGCATAAA	ACATAAAGGA	GTTTTGTTTA	585360
P	ATGTTTCGAA	TGTTTTTAT	TTAGAGCGAA	CCAGTCAAAA	GTTAAAAGAA	GATCTTTTAA	585420
P	AAAAATAGA	AGAGGTTAGT	GCAAATGGTA	TAAAAATTAA	TTTGGAACAA	AATTAATGGT	585480
P	\GAGTTTAAA	AACATAGTTA	AGTATTTTCC	AGATATTGAC	AAGCCTATTT	TGGATAGTAT	585540
7	AAATTTAAA	ATTGGGGAAG	ТТААААТТТ	TACAGTAGTT	GGTAAAAATG	GAGAAGGAAA	585600
C	SAGCACTCTA	GCCAAGATTA	TTGCCGGACT	TATTGAATTT	GATGAGGGTG	AAATATTAGT	585660
7	AATGGCATT	AAGCAAAAAA	ATTGGAATGT	AGATAAAGCT	AAAAATAATG	GTATTTATCT	585720
7	PGTTTCTCAA	GTTCCTAATT	TGAAAATGAA	TTTAAGAGTA	TGGGAATATT	TGAGTATCTA	585780
7	TTGGTTTGGT	TATGAATTTT	TCATGCCGAT	GAATAAATCT	AAGACCTACA	AATATTATAG	585840
I	ATGGCTTATG	CAATTTTATA	AAATTTCTTT	TGATTTAGAT	AAGAAAATTA	AAGATTTAAA	585900
7	TATTAAAGAG	ATTTATTTT	TACTTATTAT	TGCTGCTCTT	AAAGAGAATG	САААААТААТ	585960
7	TATTTTTGAT	GAGAGTGCTG	CTTATTTTTC	TCAAAAAGAA	GCACAAGCTT	TTATAAAATT	586020
C	CTTGTATTG	CTTAAGAAAT	CGGGAGTTGC	GTCTCTTTTT	ATTACCCACA	GCGAGATTAC	586080
2	AGATGCTATA	AAATTTAGCG	ATGAGTTTAT	TATTTTAAAA	GATGGAAAGT	GTTTTAGAAC	586140
1	AGTAAACAAA	GAATCAATTT	TGAGCAAGCT	TGAATCCTCT	AGTGACAAAG	TATTTGTTGC	586200
2	TAATTATAAA	TGCAACAAAT	TTGAAAAAGA	ТССТАТТААА	TTTAATTTGT	TTTTTGAAGA	586260
7	FTTTTGGAAG	TATGATGTTA	GTTTTTCTTT	AAATAAAAGG	GGTGTTTTAG	GGATAATTGG	586320
(CGAAGAAGCT	GTAATTAAAA	CTTGGGAAAA	ATTATTCTTA	GGAGAGCTTC	TTTTTGTTGG	586380
(GTGCATAAAA	ATTGATGGCA	TTAGATATGA	GCGAATAAAT	ATTTTTGAGT	GTAAAGCGGG	586440
Ī	ATTTTTACCC	TTAGGTATTG	GTAATTTATT	CCCCGATAAT	AGCAGCATAT	TAGATAATTT	586500
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ž	AGAGATGTAT	ATTGCAAAAA	GTTTTTTAAT	TTGTTTTTCT	CCTTTGAGCA	ATTTAGATCA	586740
(CAAAGCTTAT	AATGAAATGT	CTGTTGCTAT	TCGTAATTAT	TCAAAAGAAA	AGCCAGTTCT	586800



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					TGGCAATGAA	586860
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TGCATTCTTT A	ATATTGCATT	TTCAAATCCT	TTATTAATTT	TTCCAATTTC	TTTTTTGATA	588540
GTTTTTTTG A	AATATCAATT	TTTTCGCACT	CAAGAGTATG	ТАААТТСТТА	TTTTTCTCTT	588600



TCTTTTCAAT	TTTATGTAGC	ААТААТААТА	AATATATTGG	TTTCGTTGAT	TAAGAGAAAA	588660
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GGCCATTCCA	AAAAGTGAAG	CTGTTTTTAA	ТТСТАСТААА	GAGATGTATT	СТТТААТАСТ	589860
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TATTCGTTTT	CCGCCCCTAT	TAACAATTTC	AATTGCTGGT	GCTTTAATAT	AATCAAGGGT	590280
TTCCTTTTTT	ATTTTAAAAG	TAAATTTTAA	ATCGTTATCT	TTGAATAAAT	TTAGAAAATT	590340

P \$98/12764

AGTTGTTGAA AAGATTTTAT TAATATTTTT TTCAATATTT TTTAAAAATA GTTTATTTTG 590400 CATAATAAGA ATTATAATGA AAAAGTATAA TAAAGTGTAT TAAAGGGATT GATGTGTATC 590460 GCTTGGATGA TGAATATTCT AAAAAAGCCA AAAGAGAAGG ATATTTGGCA AGGTCTGTAT 590520 ATAAGTTGAT AGAAATTAAT GAAAAATTTT CTTTATTTTC TTCTGGCAAT GTTTTAGATA 590580 TTGGCGCATC ACCTGGCAGC TTTTCTCAGT ATGCTTATAA AAAGCTTAAA AGAGGAATTC 590640 TAGTATCTGT TGATATTAAT GATATTGGCC TTAGATATGA TGATAATTTT TATTTTATAA 590700 AGGGAGATAT CTTTTTAGAT GATACAGTTT TTAAAATTAA TACGTTTAAA CCTTATAGTC 590760 590820 TTGTAATTAG TGATGTGGCT CCCAAGACTA CTGGAAATAG ACTTGTAGAT ACCAGCAATT CTTTTAATTT AAGCATGAGA ATAATAGATT TATCACTTGA AGTTTTACTT AAAAAAGGGA 590880 ATTTACTTGT TAAAGTTTTT CAGGGAGGAG ACGAGATGCA AATTTTTAAA AAGTTTGAAA 590940 AATATTTAA ATTTGTAAAA AAAATTAGAC CCAAAGCTGT AAGGAAAAAT TCTTTTGAAA 591000 TTTATTTTT AGGCAAAAGT TTTGGCAAGT AGCAAATTAA TCAAATTGTT ATAAACAGAT 591060 TTAAAGGTAT AAAATATGTT TAGAAAAGAA AGTTCTAAAG ACAGCAGATC ACAGCTTCAA 591120 GTTGCAGGTT TTAAAATAGG AAAAGAAAGC TATGGGGTGT CAATAGAGCA CATTAGAGAA 591180 ATTATTAAAG TTCCATCAGA AGGAGTTTAT GCTATACCAA ATGTTCCCGA ATATATTATA 591240 GGTATTTATA ATCTTAGAGG CAGTATTATT CCTTTAATTA ATTTAAATAT TAAATTTGGA 591300 591360 GTTCCTTCTA TTTCGGTAAC AGAAGAAGAC ATGCTTTTAA CAGGATACTT AATAGTTAAG ATTAAAAATA AGCTTTTAGG CATTTTTGTT GATAGAGTTC TTAAAGTTAT TAGCTTTGAT 591420 GATTCTAGGG TTCAAGAACC TCCCGCTACT TTACAAACTT TAGATAGAAA ATATATATCT 591480 GGAGTTGTAA AGCTTGACGA GGCTGATAAT CTTGAGAGTG AATACTTAGT ATTAATTGAT 591540 ATAGCAAAAA TTTTTGATAA ATGCGAATTT GACGACATTC CCTATAAAGA TCAATATGAA 591600 GAATAAAGTT CTTCTTTGCA TTAATACTTT AAAGTCGGGA GCTAGTATTT TAGGCAATGA 591660 TGTTAAAGTT TATTTAGAAA CCAAGTATTT TGTTGAAGTA GTGTTAATAG ATGTTGGCAG 591720 591780 CACAGTTCTT TTGGCTGTTA ATTTGCTTCT TGAAAATGAA AACATTGATA TTCCAATTAT 591840 TTCAATTAAT ATGGGCAATG TGGGATTTTT AGCAGATATT AAGATTGAAG ATTTTAAAAA 591900 AGTCATAGAT AGATTTTTTA ACAATTCTTT GGTTATTAAT AAAAAATTTT TGCTTCATGT 591960 592020 AACAGTTTCT CAACACGGTA AAGATTTAAT TTCTAAATAT GCTTTAAACG ATATTATTAT TCGCTCAAGC GTTCTTAATA AAATGATTTA TGTAGATCTT ATGGTTAATT CTGAGAGTTT 592080 TTTATCATAC AAAAGTGATG GGATAATTGT GTCTACTCCA ACAGGCTCAA CAGGATATTC 592140





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TTTCTCAGCA GGGGGTCCTA TTTTAGAAGC AGATCTTGAG GGATTTTYAC TTACGCCTAT 592200 TTCTCCACAT TCTGTTTATA ATCGTTCTTT TGTGTTCTCT AAATTAAGTA AACTTTCCAT 592260 TTCTTTTCA AAGGAATATT TTATAGCAGC AGCATCAATT TTTTTAGATG GAATTAATTT 592320 TGGTTCTTTC GGAGTTGACG TTGTTTTTGA ATTTAAAATT TCTTCTCAAA GCTTGAATTT 592380 TGTTTCATTT TGTACGGATA CTTTTGTTAA GAGATTAAAA AACAAATTAT TGTAAGTTCA 592440 ATGTTTTTT AAACAGTGTT CTTTTTAATA AAAATTTTTC TTGGTTGTTG TTTGAATGTT 592500 TGGTTTAATC TAATCTTTAA TAAGGTTAGA GGATTTTACA TGTTTTATTA TAAGGATTTT 592560 AATGTTTTGT TTATTCGCAG AGTTAATGTT TAAAGGGTGT TTTTTTATAAA TTTATAACTT 592620 GTTAAAATAT TTAATATAT TTTTGCTAAA ATCAATATAT TAGATAGCTG ATAAATCACC 592680 CTTTTATGAG TTTTAGGCAG GTGTTTATGG ATTTAATTGA TAATGAAAAAT TATAAAAAAA 592740 TAGTGTACAT TAATAATCTT GTTTTAAGGA CTTTAAATGA TATAGCAGCT ATAAAAGAGA 592800 CTGGCGAATT TACATCAAAT GCTAAACTTT CATTTAATCT TATTGATTTC AATTTAAATG 592860 TTTTAAGTTA TATTTCTTCT TTAAATTATT TTTATACTAG GCCTAGATTG AAAGTAAATT 592920 592980 TTCTTAGTAT TACTCCAAGA GAGTTGATTG AAATGCCTCA GGCTCTTAAT TTAAATCCAG 593040 AAGAGAGGTT TTTAATTATT AAAAAATTAG GTTATTTAAT TGATTTGGCT AAAATTTTTA 593100 GCAAAAAGA TTCTAAAACA CTTGTTTTC TTGAGGATAT GTATCTCAAA TTTATTGTTT 593160 TTTCTAAAAA TATTATTGAT TTTAGAGATT TTTCTAAGAA TTTAAAAACTT GAGAGTCCTT 593220 ATTATAAATT TCAATTTGAA CACCTTATTA AAGTGTTGGA GCTTTTAGAA GAAGGAGCTT 593280 TTATTTTAAG GGGCAAATAT GAGATTAGTG GATCTCATGA ATTTGGACTG CATTCTCTTG 593340 GTTATCTTGA AGCTGGAAGA GCTTTGGCTA CTATAGCCTC TCAAAAAGAA GCTGCTGAAA 593400 AATTTTCAAG GTTTCATGGA GTTTGGTCTT CAAAGTTTAG TTCAGATTTA ATTAAAGTAA 593460 AATAGATAAA TTAAGGTGGG GAGAAAGTAG TTATATTGAG TTTTAATGTA GAAGAGGGCA 593520 CTATTAAATT CAAAAAATTA AAATTTTTTT TGATTCTAAG CTTGTTTTTA TTATTATAA 593580 TTTTGATTGA TTTTTTTATA AGATCTACTA TGAATGTATC TAATTTTTAT GATTTTAAGA 593640 ATTTTGAAAA TAAATCTGAT TGTAAAAATA TAAATTTAAG TAAGAATGTT TTTGTATCAA 593700 ATAAGGTTTT AAGTCTTAAT TTCGGGGAAT CTTGTTATTC TCTTTTAAGT GATAACTTAA 593760 TAAGTTATTC AGACTATTAT TATGTGCTTT TTAATTCCGG CGAAGATTAT TCTGTTTTTT 593820 CTGTTAAAAA CAATAAATTT TTATTTACAC TCAAGCTAAA GGATTTTGTT TTTGCAATAA



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495	
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GAAGTAATTA	TTTATGATGA	AATTGCTTTA	TATTGATAAT	TTGAAATTTT	TAAAAGGCAA	604140
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AGACATTGAG	TCTTATTTTT	ТААААТСАТТ	TAGAAGATTG	TTTAAGTTGC	CCGATCTAAA	604260
ATTAGTAGAA	TTACAAGAAA	AAGTTATTCA	AAGGACCAAA	GCCAAAGTTG	CTATTCTAGG	604320
GTCTAAGTCT	ГАТССТААТА	AGCTTAAAAG	AATTTATGAC	CCTCCTTTTG	СТАТТТАСТА	604380
CAAGGGCAAT	TTACCAGATT	GTTCTTTATT	ATCTTGGGCT	GTTGTTGGTT	СТАGAAAAAT	604440
TAGTAAAACT (CTTGCTGAGA	GAACAAGGGA	ATTTTCTTCA	CATCTTGCAA .	AGAATGGTGT	604500





			499			
AGAGATTATT	TCTGGATTTG	CAATTGGGGC	AGATATTGAG	GCTCATATAG	CAGCAATAAA	604560
TGAGAATAAG	AGAACATTTG	CTGTTATTCC	AACAGATATT	GACAATATTT	ATCCTAGGCA	604620
AAATCGAAAA	TATGTTTCCA	AGCTTTTAGA	ACAAGGTGGA	GGAATAATTA	CTGAGACTTT	604680
GCCATTTGAT	AAAATTCAAA	ATTATTTTTT	TGCCAAAAGA	AATAGATTGG	TATCAGGTCT	604740
GTCTGATGCT	ATTTTTATAA	CATATGCACC	CTTGAAATCA	GGAGCTTTAA	TTACAGCTGA	604800
GCTTGGTCTT	GACTTAGGAC	TTGATGTTTA	TGTTTATGAT	TTAGATTTTT	GTGGTGATGG	604860
AGCTGTAAAA	TTGCATGATT	TTGGTGCGCA	AGAAATAAAA	ACCGTTAAGG	ATCTTTATGC	604920
ТТТАТТААТТ	ATTAAATATG	TAGATTCCAA	TAATATTGAA	GATGATTCTA	AAGAGTGTTG	604980
TAATTGTAAA	AATGTATCTG	ATGTTCTTAT	TGGGGAACTT	TTAAAAGAGG	TATGTAAATA	605040
GGGGGGTAAT	ATGAGCTTTA	AAGGAACCAC	AGTTATTGCA	ATAAAAAAAA	ATGGTAAGAC	605100
TGTGGTGGCA	GCAGATGGAC	AAGTAACTTT	TGGACATACT	GTTTTAAAGA	GTAATGCTAT	605160
TAAAATACGA	AAATTGCTTA	ATGGGAAAAT	TTTGGCAGGA	TTTGCAGGTT	CAACATCTGA	605220
TGCAATTACT	CTTTTTGAAA	AATTTGAAGA	АААААТСААА	GCAAAAGGTG	ATGGCTTGAT	605280
TGACATTAAA	AGGGCGGCTG	TTGACCTTGC	AAAAGATTGG	CGTTCTGACA	AAATACTGCA	605340
TAAGCTTGAG	GCTATGATGC	TTGTTGCTGA	TTCTAACAAT	ATTCTTTTGA	TTTCTGGTAC	605400
TGGTGATGTT	GTTGAGCCTG	AAGAGGATGT	TATTTCGATT	GGCAGTGGTG	GTAATTATGC	605460
ATATTCAGCA	GCTCTTGCTT	ACATGGAGAA	СААААААТТА	AGCGCTTTTG	AGGTTGCACT	605520
TAGATCTTTA	AAAATAGCAG	CAAGAGTGTG	TATATATACT	AATTCTAATA	TTGTGCTTGA	605580
GGAGATTGAA	AATGAATAAA	TTAGAAGAGC	ACTATATAGT	TCCCAAAGAT	GTAGTTGCAG	605640
AACTTGATAA	АТАТАТААТА	GGTCAAGACG	AAGCTAAAAA	ATTAGTATCA	ATTGCTCTTG	605700
TTAATAGATA	TATAAGGTCT	AGGCTTCCAA	AAGAAATAAA	AGATGAGGTA	ATGCCTAAAA	605760
ACATTATTAT	GATTGGATCA	ACTGGCATTG	GGAAGACCGA	GATTGCAAGA	AGACTTTCTA	605820
AATTAATTAA	AGCTCCTTTT	ATTAAAGTTG	AGGCTACAAA	ATATACTGAG	GTTGGTTATG	605880
TTGGTCGTGA	TGTTGAATCT	ATGGTTAGAG	ATTTAATGAG	CATTGCAGTT	AATATGGTAA	605940
AAGAAGAGAT	GTATAGTACT	GTAAGAGATG	ATGCTTTAGT	AAGAACAGAG	GAGAGAATAG	606000
TTGATAGTCT	TTTTAAGGGA	TCTAGTAATT	CTGAGAATAT	GGATCCAAAT	GAAATAAAGG	606060
CGGAAGAAAA	GGTAAAAGAG	AAGCTTAGAA	AAAAGCTTAG	AGCAGGTGAG	CTTGATGATA	606120
CTACTATTGA	AATACAAATT	TCTAGTAAAA	TGCCATTTTC	TACAATAGAA	ATATTTACGG	606180
GTGGTAATTT	TĠAAGAGATT	GATATGGGAA	TTGGCGGTTT	GCTGGGTAAT	ATATTTGATA	606240
GAAAAAAGAA	AAGAGAATTG	AAGATTAAAA	AAGCAAAGGA	ATTATTA	GCAGAAGAGC	606300